



## Clinical trial results:

**An open-label, multicenter rollover study to provide continued treatment with anetumab ravtansine for participants with solid tumors who were enrolled in previous Bayer-sponsored studies**

### Summary

EudraCT number	2019-000061-20
Trial protocol	FR BE PL IT
Global end of trial date	18 May 2022

### Results information

Result version number	v1
This version publication date	14 May 2023
First version publication date	14 May 2023

### Trial information

#### Trial identification

Sponsor protocol code	20322
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#### Additional study identifiers

ISRCTN number	-
ClinicalTrials.gov id (NCT number)	NCT03926143
WHO universal trial number (UTN)	-

Notes:

#### Sponsors

Sponsor organisation name	Bayer AG
Sponsor organisation address	Kaiser-Wilhelm-Allee, Leverkusen, Germany, D-51368
Public contact	Therapeutic Area Head, Bayer AG, +49 30 300139003, clinical-trials-contact@bayer.com
Scientific contact	Therapeutic Area Head, Bayer AG, +49 30 300139003, clinical-trials-contact@bayer.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	18 May 2022
Is this the analysis of the primary completion data?	Yes
Primary completion date	18 May 2022
Global end of trial reached?	Yes
Global end of trial date	18 May 2022
Was the trial ended prematurely?	Yes

Notes:

## General information about the trial

Main objective of the trial:

This study was a rollover study to permit subjects who received anetumab ravtansine in an applicable Bayer sponsored anetumab ravtansine parent study to continue treatment or follow-up at the time of parent study closure.

Main objective is safety.

Protection of trial subjects:

The conduct of this clinical study met all local legal and regulatory requirements. The study was conducted in accordance with ethical principles that have their origin in the Declaration of Helsinki and the International Council for Harmonization guideline E6: Good Clinical Practice. Before entering the study, the informed consent was read by and explained to all the subjects. Participating subjects signed informed consent form and could withdraw from the study at any time without any disadvantage and without having to provide a reason for this decision. Only investigators qualified by training and experience were selected as appropriate experts to investigate the study drug.

Background therapy: -

Evidence for comparator: -

Actual start date of recruitment	03 June 2019
Long term follow-up planned	Yes
Long term follow-up rationale	Safety, Efficacy
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	France: 1
Country: Number of subjects enrolled	Italy: 1
Country: Number of subjects enrolled	Poland: 2
Country: Number of subjects enrolled	United States: 6
Worldwide total number of subjects	10
EEA total number of subjects	4

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)	6
From 65 to 84 years	4
85 years and over	0

## Subject disposition

### Recruitment

Recruitment details:

The study was conducted at 7 study centers in 4 countries worldwide between 03-Jun-2019 (first subject first visit) and 18-May-2022 (last subject last visit).

### Pre-assignment

Screening details:

A total of 10 subjects were screened in this study; of whom 9 subjects started study treatment and 1 subject was a screening failure. Entering subjects had to have been treated with anetumab ravtansine in an applicable Bayer sponsored anetumab ravtansine parent study.

### Period 1

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

### Arms

<b>Arm title</b>	Anetumab ravtansine
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Arm description:

Adult subjects with solid tumors who received anetumab-ravtansine treatment as monotherapy, or in combination with gemcitabine in an applicable Bayer-sponsored anetumab ravtansine study.

8 subjects received anetumab ravtansine monotherapy and 1 subject received anetumab ravtansine in combination with gemcitabine. Pooled results were reported.

Arm type	Experimental
Investigational medicinal product name	Gemcitabine
Investigational medicinal product code	
Other name	
Pharmaceutical forms	Infusion
Routes of administration	Intravenous use

Dosage and administration details:

Gemcitabine as per the dosing instructions from the parent study protocol.

Investigational medicinal product name	Anetumab ravtansine
Investigational medicinal product code	BAY94-9343
Other name	
Pharmaceutical forms	Infusion
Routes of administration	Intravenous use

Dosage and administration details:

Anetumab ravtansine as per the dosing instructions from the parent study protocol in an every 3 weeks (Q3W) schedule.

<b>Number of subjects in period 1<sup>[1]</sup></b>	Anetumab ravtansine
Started	9
Completed	0
Not completed	9
Physician decision	3
Adverse event, non-fatal	2

Subject decision: Covid-19 Pandemic related	1
Progressive disease	3

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Notes:

[1] - The number of subjects reported to be in the baseline period are not the same as the worldwide number enrolled in the trial. It is expected that these numbers will be the same.

Justification: In total, 10 subjects were enrolled and 9 subjects started the treatment.

## Baseline characteristics

### Reporting groups

Reporting group title	Anetumab ravtansine
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Reporting group description:

Adult subjects with solid tumors who received anetumab-ravtansine treatment as monotherapy, or in combination with gemcitabine in an applicable Bayer-sponsored anetumab ravtansine study.

8 subjects received anetumab ravtansine monotherapy and 1 subject received anetumab ravtansine in combination with gemcitabine. Pooled results were reported.

Reporting group values	Anetumab ravtansine	Total	
Number of subjects	9	9	
Age Categorical Units: Subjects			
Age Continuous Units: years arithmetic mean standard deviation	50.7 ± 16.2	-	
Gender Categorical Units: Subjects			
Female	3	3	
Male	6	6	
Race Units: Subjects			
White	6	6	
Asian	2	2	
Not reported	1	1	
Ethnicity Units: Subjects			
Unknown or Not Reported	9	9	

## End points

### End points reporting groups

Reporting group title	Anetumab ravtansine
Reporting group description: Adult subjects with solid tumors who received anetumab-ravtansine treatment as monotherapy, or in combination with gemcitabine in an applicable Bayer-sponsored anetumab ravtansine study. 8 subjects received anetumab ravtansine monotherapy and 1 subject received anetumab ravtansine in combination with gemcitabine. Pooled results were reported.	

### Primary: Number of subjects with TEAEs, TESAEs and Drug-related TEAEs and TESAEs

End point title	Number of subjects with TEAEs, TESAEs and Drug-related TEAEs and TESAEs <sup>[1]</sup>
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End point description:

Treatment emergent adverse events (TEAEs) were defined as AEs starting or worsening during the treatment period. The treatment period extended from the first date of study treatment in this study until the safety follow-up (30 days after the last administration of study treatment). TESAEs: Treatment emergent serious adverse events.

End point type	Primary
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End point timeframe:

Approximately 3 years (from first study treatment until safety follow-up)

Notes:

[1] - No statistical analyses have been specified for this primary end point. It is expected there is at least one statistical analysis for each primary end point.

Justification: As study was small and incoming population was heterogeneous and subject to selection bias, no inferential statistical analysis was performed.

End point values	Anetumab ravtansine			
Subject group type	Reporting group			
Number of subjects analysed	9			
Units: Subjects				
Any TEAE	9			
Serious TEAE	2			
Any study drug-related TEAE	8			
Any study drug-related Serious TEAE	0			

### Statistical analyses

No statistical analyses for this end point

### Secondary: Overall survival

End point title	Overall survival
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End point description:

Overall survival (OS) defined as the time from first treatment in this study until death from any cause. Data on survival were collected by the site. Time frame was reduced due to early termination of the study. Table reports Kaplan-Meier median with Brookmeyer-Crowley confidence intervals. 99999

indicates value cannot be estimated due to censored data.

End point type	Secondary
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End point timeframe:

Approximately 3 years (from first study treatment until safety follow-up)

<b>End point values</b>	Anetumab ravtansine			
Subject group type	Reporting group			
Number of subjects analysed	9 <sup>[2]</sup>			
Units: Months				
median (confidence interval 95%)				
25th percentile	17.6 (6.9 to 34.1)			
Median	34.1 (6.9 to 99999)			
75th percentile	99999 (28.5 to 99999)			

Notes:

[2] - Number of subjects: with event (5) and censored (4)

### **Statistical analyses**

No statistical analyses for this end point

## Adverse events

### Adverse events information

Timeframe for reporting adverse events:

For TEAE: After the first study intervention up to 30 days after the end of study intervention, approximately 2 years. For the deaths (all causes) considers all deaths that occurred at any time during the study before the last contact.

Assessment type	Non-systematic
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### Dictionary used

Dictionary name	MedDRA
Dictionary version	25.0

### Reporting groups

Reporting group title	Anetumab ravtansine
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Reporting group description:

Adult patients with solid tumors who received anetumab-ravtansine treatment as monotherapy, or in combination with gemcitabine in an applicable Bayer-sponsored anetumab ravtansine study. 8 subjects received anetumab ravtansine monotherapy and 1 subject received anetumab ravtansine in combination with gemcitabine. Pooled results were reported.

<b>Serious adverse events</b>	Anetumab ravtansine		
Total subjects affected by serious adverse events			
subjects affected / exposed	2 / 9 (22.22%)		
number of deaths (all causes)	5		
number of deaths resulting from adverse events	0		
Cardiac disorders			
Restrictive cardiomyopathy			
subjects affected / exposed	1 / 9 (11.11%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
General disorders and administration site conditions			
Oedema peripheral			
subjects affected / exposed	1 / 9 (11.11%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Respiratory, thoracic and mediastinal disorders			
Dyspnoea			
subjects affected / exposed	1 / 9 (11.11%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		

Infections and infestations			
Urosepsis			
subjects affected / exposed	1 / 9 (11.11%)		
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 %

<b>Non-serious adverse events</b>	Anetumab ravtansine		
Total subjects affected by non-serious adverse events			
subjects affected / exposed	9 / 9 (100.00%)		
Neoplasms benign, malignant and unspecified (incl cysts and polyps)			
Transitional cell carcinoma			
subjects affected / exposed	1 / 9 (11.11%)		
occurrences (all)	1		
Vascular disorders			
Hypotension			
subjects affected / exposed	1 / 9 (11.11%)		
occurrences (all)	1		
General disorders and administration site conditions			
Fatigue			
subjects affected / exposed	1 / 9 (11.11%)		
occurrences (all)	1		
Oedema peripheral			
subjects affected / exposed	2 / 9 (22.22%)		
occurrences (all)	3		
Chest pain			
subjects affected / exposed	1 / 9 (11.11%)		
occurrences (all)	1		
Asthenia			
subjects affected / exposed	2 / 9 (22.22%)		
occurrences (all)	2		
Reproductive system and breast disorders			
Gynaecomastia			

subjects affected / exposed occurrences (all)	2 / 9 (22.22%) 2		
Respiratory, thoracic and mediastinal disorders			
Upper-airway cough syndrome subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 1		
Oropharyngeal pain subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 1		
Pleural effusion subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 1		
Nasal congestion subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 1		
Epistaxis subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 1		
Dyspnoea subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 2		
Investigations			
Activated partial thromboplastin time prolonged subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 1		
Blood alkaline phosphatase increased subjects affected / exposed occurrences (all)	2 / 9 (22.22%) 6		
Schirmer's test abnormal subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 1		
Transaminases increased subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 1		
Weight decreased			

subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 4		
Neutrophil count decreased subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 2		
Lipase increased subjects affected / exposed occurrences (all)	2 / 9 (22.22%) 3		
Blood thyroid stimulating hormone increased subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 1		
Blood creatinine increased subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 2		
Blood bilirubin increased subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 1		
Aspartate aminotransferase increased subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 3		
Amylase increased subjects affected / exposed occurrences (all)	2 / 9 (22.22%) 3		
Injury, poisoning and procedural complications			
Corneal abrasion subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 1		
Foot fracture subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 1		
Cardiac disorders			
Restrictive cardiomyopathy subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 2		
Pericardial effusion			

<p>subjects affected / exposed occurrences (all)</p> <p>Sinus tachycardia subjects affected / exposed occurrences (all)</p>	<p>1 / 9 (11.11%) 1</p> <p>1 / 9 (11.11%) 1</p>		
<p>Nervous system disorders</p> <p>Peripheral motor neuropathy subjects affected / exposed occurrences (all)</p> <p>Neuropathy peripheral subjects affected / exposed occurrences (all)</p>	<p>2 / 9 (22.22%) 6</p> <p>1 / 9 (11.11%) 1</p>		
<p>Blood and lymphatic system disorders</p> <p>Thrombocytopenia subjects affected / exposed occurrences (all)</p> <p>Neutropenia subjects affected / exposed occurrences (all)</p> <p>Lymphopenia subjects affected / exposed occurrences (all)</p> <p>Anaemia subjects affected / exposed occurrences (all)</p> <p>Leukopenia subjects affected / exposed occurrences (all)</p>	<p>1 / 9 (11.11%) 3</p> <p>1 / 9 (11.11%) 3</p> <p>1 / 9 (11.11%) 1</p> <p>5 / 9 (55.56%) 10</p> <p>1 / 9 (11.11%) 1</p>		
<p>Eye disorders</p> <p>Keratitis subjects affected / exposed occurrences (all)</p> <p>Cataract subjects affected / exposed occurrences (all)</p> <p>Corneal epithelial microcysts</p>	<p>1 / 9 (11.11%) 1</p> <p>1 / 9 (11.11%) 1</p>		

subjects affected / exposed occurrences (all)	2 / 9 (22.22%) 2		
Corneal disorder subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 1		
Vision blurred subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 1		
Gastrointestinal disorders			
Toothache subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 1		
Nausea subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 1		
Diarrhoea subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 2		
Constipation subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 1		
Abdominal pain subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 2		
Abdominal discomfort subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 1		
Endocrine disorders			
Hyperthyroidism subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 1		
Musculoskeletal and connective tissue disorders			
Pain in extremity subjects affected / exposed occurrences (all)	1 / 9 (11.11%) 1		
Muscle spasms			

<p>subjects affected / exposed occurrences (all)</p> <p>Back pain subjects affected / exposed occurrences (all)</p> <p>Arthralgia subjects affected / exposed occurrences (all)</p>	<p>1 / 9 (11.11%) 1</p> <p>1 / 9 (11.11%) 1</p> <p>2 / 9 (22.22%) 2</p>		
<p>Infections and infestations</p> <p>Conjunctivitis subjects affected / exposed occurrences (all)</p> <p>Sinusitis subjects affected / exposed occurrences (all)</p> <p>Skin infection subjects affected / exposed occurrences (all)</p>	<p>1 / 9 (11.11%) 2</p> <p>1 / 9 (11.11%) 1</p> <p>1 / 9 (11.11%) 1</p>		
<p>Metabolism and nutrition disorders</p> <p>Hyperglycaemia subjects affected / exposed occurrences (all)</p> <p>Cachexia subjects affected / exposed occurrences (all)</p> <p>Hypokalaemia subjects affected / exposed occurrences (all)</p>	<p>2 / 9 (22.22%) 2</p> <p>1 / 9 (11.11%) 1</p> <p>1 / 9 (11.11%) 1</p>		

## More information

### Substantial protocol amendments (globally)

Were there any global substantial amendments to the protocol? No

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### 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.

Due to the small sample size and heterogeneous population, survival distributions and the extent of long-term survival in the applicable populations cannot be reliably estimated from the study results.

Notes: