



## Clinical trial results:

### Phase 3, Randomized, Double-Blind Trial of Pegylated Liposomal Doxorubicin (PLD) Plus AMG 386 or Placebo in Women With Recurrent Partially Platinum Sensitive or Resistant Epithelial Ovarian, Primary Peritoneal, or Fallopian Tube Cancer

#### Summary

EudraCT number	2009-017946-30
Trial protocol	HU GB SK AT BE IT DK DE PL GR LV ES FR
Global end of trial date	19 April 2017

#### Results information

Result version number	v1 (current)
This version publication date	02 May 2018
First version publication date	02 May 2018

#### Trial information

##### Trial identification

Sponsor protocol code	20060517
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##### Additional study identifiers

ISRCTN number	-
ClinicalTrials.gov id (NCT number)	NCT01281254
WHO universal trial number (UTN)	-

Notes:

#### Sponsors

Sponsor organisation name	Amgen Inc.
Sponsor organisation address	One Amgen center Drive, Thousand Oaks, CA, United States, 91320
Public contact	IHQ Medical Info-Clinical Trials, Amgen (EUROPE) GmbH, MedInfoInternational@amgen.com
Scientific contact	IHQ Medical Info-Clinical Trials, Amgen (EUROPE) GmbH, MedInfoInternational@amgen.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	19 April 2017
Is this the analysis of the primary completion data?	No
Global end of trial reached?	Yes
Global end of trial date	19 April 2017
Was the trial ended prematurely?	No

Notes:

## General information about the trial

Main objective of the trial:

The primary objective was to determine if trebananib plus pegylated liposomal doxorubicin (PLD) is superior to placebo plus PLD as measured by progression-free survival (PFS), defined as the time from randomization to the earliest of the dates of first radiologic disease progression per Response Evaluation Criteria in Solid Tumors 1.1 with modifications (RECIST 1.1 mod) or death from any cause in subjects with recurrent partially platinum sensitive or resistant epithelial ovarian, primary peritoneal or fallopian tube cancer.

Protection of trial subjects:

This study was conducted in accordance with International Conference on Harmonisation (ICH) Good Clinical Practice (GCP) regulations/guidelines, United States Food and Drug Administration (FDA) regulations/guidelines, and country-specific national and local laws.

A copy of the protocol, proposed informed consent form (ICF), other written subject information, and any proposed advertising material was submitted to the Independent Ethics Committee (IEC) or Institutional Review Board (IRB) for written approval. A copy of the IEC/IRB approval was received by the sponsor before recruitment of subjects into the study.

All subjects provided written informed consent before undergoing any study-related procedures, including screening procedures.

Background therapy: -

Evidence for comparator: -

Actual start date of recruitment	18 April 2011
Long term follow-up planned	Yes
Long term follow-up rationale	Safety, Efficacy
Long term follow-up duration	60 Months
Independent data monitoring committee (IDMC) involvement?	Yes

Notes:

## Population of trial subjects

### Subjects enrolled per country

Country: Number of subjects enrolled	Canada: 9
Country: Number of subjects enrolled	United States: 23
Country: Number of subjects enrolled	Australia: 11
Country: Number of subjects enrolled	Austria: 15
Country: Number of subjects enrolled	Belgium: 37
Country: Number of subjects enrolled	Denmark: 2
Country: Number of subjects enrolled	France: 4
Country: Number of subjects enrolled	Germany: 12
Country: Number of subjects enrolled	Italy: 37
Country: Number of subjects enrolled	New Zealand: 7
Country: Number of subjects enrolled	United Kingdom: 26

Country: Number of subjects enrolled	Hong Kong: 9
Country: Number of subjects enrolled	Hungary: 14
Country: Number of subjects enrolled	Poland: 3
Country: Number of subjects enrolled	Singapore: 5
Country: Number of subjects enrolled	Slovakia: 9
Worldwide total number of subjects	223
EEA total number of subjects	159

Notes:

<b>Subjects enrolled per age group</b>	
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)	147
From 65 to 84 years	76
85 years and over	0

## Subject disposition

### Recruitment

Recruitment details:

This study was conducted at 69 centers worldwide. Participants were enrolled from 18 April 2011 to 12 November 2013. Enrollment was put on hold from 23 November 2011 until 10 January 2013 because of a global shortage of PLD. On 23 October 2013, Amgen closed the study to subject screening because of a further imminent shortage of PLD.

### Pre-assignment

Screening details:

Participants were randomized in a 1:1 ratio to 1 of 2 treatment groups. Randomization was stratified based on platinum-free interval (PFI) status (PFI  $\geq$  0 months and  $\leq$  6 months versus PFI > 6 months and  $\leq$  12 months), presence / absence of measurable disease, and region (North America, Western Europe/Australia, Rest of World [ROW]).

### Period 1

Period 1 title	Overall Study (overall period)
Is this the baseline period?	Yes
Allocation method	Randomised - controlled
Blinding used	Double blind
Roles blinded	Investigator, Subject

### Arms

Are arms mutually exclusive?	Yes
<b>Arm title</b>	Placebo + PLD

Arm description:

Participants received pegylated liposomal doxorubicin (PLD) 50 mg/m<sup>2</sup> every 4 weeks (Q4W) plus placebo to trebananib administered intravenously (IV) every week (QW) until disease progression per RECIST 1.1 mod, unacceptable toxicity, withdrawal of consent, or death.

Arm type	Placebo
Investigational medicinal product name	Placebo to Trebananib
Investigational medicinal product code	
Other name	
Pharmaceutical forms	Powder for solution for infusion
Routes of administration	Intravenous use

Dosage and administration details:

Administered as an IV infusion

<b>Arm title</b>	Trebananib + PLD
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Arm description:

Participants received PLD 50 mg/m<sup>2</sup> every 4 weeks (Q4W) plus trebananib 15 mg/kg administered intravenously (IV) every week (QW) until disease progression per RECIST 1.1 mod, unacceptable toxicity, withdrawal of consent, or death.

Arm type	Experimental
Investigational medicinal product name	Trebananib
Investigational medicinal product code	AMG 386
Other name	
Pharmaceutical forms	Powder for solution for infusion
Routes of administration	Intravenous use

Dosage and administration details:

15 mg/kg administered as an IV infusion

<b>Number of subjects in period 1</b>	Placebo + PLD	Trebananib + PLD
Started	109	114
Received Trebananib/Placebo	108	112
Completed	0	0
Not completed	109	114
Consent withdrawn by subject	2	6
Death	94	102
Other	12	5
Lost to follow-up	1	1

## Baseline characteristics

### Reporting groups

Reporting group title	Placebo + PLD
Reporting group description:	
Participants received pegylated liposomal doxorubicin (PLD) 50 mg/m <sup>2</sup> every 4 weeks (Q4W) plus placebo to trebananib administered intravenously (IV) every week (QW) until disease progression per RECIST 1.1 mod, unacceptable toxicity, withdrawal of consent, or death.	
Reporting group title	Trebananib + PLD
Reporting group description:	
Participants received PLD 50 mg/m <sup>2</sup> every 4 weeks (Q4W) plus trebananib 15 mg/kg administered intravenously (IV) every week (QW) until disease progression per RECIST 1.1 mod, unacceptable toxicity, withdrawal of consent, or death.	

Reporting group values	Placebo + PLD	Trebananib + PLD	Total
Number of subjects	109	114	223
Age Categorical			
Units: Subjects			
Adults (18-64 years)	79	68	147
From 65-84 years	30	46	76
85 years and over	0	0	0
Age Continuous			
Units: years			
arithmetic mean	59.7	60.3	
standard deviation	± 9.2	± 9.6	-
Gender Categorical			
Units: Subjects			
Female	109	114	223
Male	0	0	0
Race			
Units: Subjects			
Asian	12	10	22
Black (or African American)	2	1	3
Native Hawaiian or Other Pacific Islander	1	0	1
White	92	102	194
Other	2	1	3
Primary Tumor Type			
Units: Subjects			
Fallopian tube cancer	1	8	9
Ovarian cancer	95	98	193
Primary peritoneal carcinoma	13	8	21
Eastern Cooperative Oncology Group (ECOG) Performance Status			
A scale to assess a patient's disease status. 0 = Fully active, able to carry out all pre-disease performance without restriction; 1 = Restricted in physically strenuous activity, ambulatory and able to carry out work of a light nature; 2 = Ambulatory and capable of all self-care, unable to carry out any work activities. Up and about > 50% of waking hours; 3 = Capable of only limited self-care, confined to bed or chair > 50% of waking hours; 4 = Completely disabled, confined to bed or chair; 5 = Dead.			
Units: Subjects			
0 (Fully active)	67	75	142
1 (Restricted but ambulatory)	41	39	80

2 (Ambulatory but unable to work)	1	0	1
Histologic Type			
Units: Subjects			
Mucinous	1	4	5
Serous	82	89	171
Endometrioid	7	6	13
Clear cell	4	4	8
Undifferentiated	6	3	9
Other	8	8	16
Not applicable	1	0	1
Histologic Grade			
Units: Subjects			
Well differentiated	5	3	8
Moderately differentiated	18	14	32
Poorly differentiated	70	77	147
Unknown	16	20	36
Number of Lines of Prior Therapy			
Units: Subjects			
One	40	45	85
Two	46	45	91
Three	23	24	47
Measurable Disease at Baseline			
Units: Subjects			
Yes	96	99	195
No	13	15	28
Region			
Units: Subjects			
North America	14	18	32
Western Europe/Australia	75	76	151
Rest of the World	20	20	40
Platinum-free Interval (PFI) Status			
Platinum-free interval (PFI) was defined as the time from the last dose of the last platinum-containing regimen until the first date of progression was noted following discontinuation of the last prior platinum-containing agent.			
Units: Subjects			
≤ 6 months	60	62	122
> 6 months to ≤ 12 months	49	52	101
Enrollment Period			
Units: Subjects			
Enrolled prior to hold	26	32	58
Enrolled after hold	83	82	165

## End points

### End points reporting groups

Reporting group title	Placebo + PLD
Reporting group description: Participants received pegylated liposomal doxorubicin (PLD) 50 mg/m <sup>2</sup> every 4 weeks (Q4W) plus placebo to trebananib administered intravenously (IV) every week (QW) until disease progression per RECIST 1.1 mod, unacceptable toxicity, withdrawal of consent, or death.	
Reporting group title	Trebananib + PLD
Reporting group description: Participants received PLD 50 mg/m <sup>2</sup> every 4 weeks (Q4W) plus trebananib 15 mg/kg administered intravenously (IV) every week (QW) until disease progression per RECIST 1.1 mod, unacceptable toxicity, withdrawal of consent, or death.	

### Primary: Progression-free Survival (PFS)

End point title	Progression-free Survival (PFS)
End point description: PFS was defined as the time from the date of randomization to the earliest of the dates of first radiologic disease progression per RECIST 1.1 with modifications, based on investigator assessment, or death from any cause. Subjects who did not meet these criteria by the analysis data cutoff date had their PFS time censored at the latest of their last evaluable radiologic disease assessment date. Events of radiographic progression per RECIST 1.1 with modifications that occurred after initiation of subsequent anticancer therapy were not considered PFS events and were censored at the last evaluable radiographic tumor assessment before the initiation of subsequent anticancer therapy. Deaths that occurred after initiation of subsequent anticancer therapy were considered PFS events.	
End point type	Primary
End point timeframe: From randomization to the data cut off-date of 19 April 2017; median follow-up time was 15.2 months (interquartile range [IQR], 8.8-25.4) in the Placebo arm and 17.3 months (IQR, 8.4-27.7) in the Trebananib group	

End point values	Placebo + PLD	Trebananib + PLD		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	109	114		
Units: months				
median (confidence interval 95%)	7.3 (4.8 to 8.8)	7.6 (7.3 to 9.2)		

### Statistical analyses

Statistical analysis title	Primary Evaluation
Statistical analysis description: A stratified log-rank test was used for the primary comparison of PFS.	
Comparison groups	Trebananib + PLD v Placebo + PLD



Number of subjects included in analysis	223
Analysis specification	Pre-specified
Analysis type	superiority
P-value	= 0.54 <sup>[1]</sup>
Method	Stratified Log-rank Test

Notes:

[1] - Stratified by PFI status and enrollment before PLD shortage

<b>Statistical analysis title</b>	Piece-wise Cox Model Analysis of PFS
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Statistical analysis description:

A stratified piecewise Cox regression model using 16-week intervals was used to estimate the PFS hazard ratio (HR) and 2-sided 95% confidence interval (CI) for trebananib in combination with PLD relative to placebo and PLD.

Comparison groups	Placebo + PLD v Trebananib + PLD
Number of subjects included in analysis	223
Analysis specification	Pre-specified
Analysis type	superiority
P-value	= 0.56 <sup>[2]</sup>
Method	Cox proportional hazards model
Parameter estimate	Overall hazard ratio
Point estimate	0.92
Confidence interval	
level	95 %
sides	2-sided
lower limit	0.69
upper limit	1.22

Notes:

[2] - Stratified Cox proportional hazards models fit with treatment as covariate, and additional time-dependent indicators for each additional time interval for the test arm. Stratification factors are PFI status and enrollment prior to PLD shortage.

## Secondary: Overall Survival

End point title	Overall Survival
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End point description:

Overall survival was defined as the time from the randomization date to the date of death from any cause. Subjects who did not die by the analysis data cutoff date were censored at their last contact date prior to the data cutoff date. Subjects known to be alive prior to the data cutoff date were censored at the last contact prior to the cutoff date. Subjects known to be alive or dead after the data cutoff date were censored at the data cutoff date.

End point type	Secondary
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End point timeframe:

From randomization to the data cut off-date of 19 April 2017; median follow-up time was 15.2 months (interquartile range [IQR], 8.8-25.4) in the Placebo arm and 17.3 months (IQR, 8.4-27.7) in the Trebananib group.

End point values	Placebo + PLD	Trebananib + PLD		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	109	114		
Units: months				
median (confidence interval 95%)	15.8 (13.5 to 20.5)	18.5 (13.4 to 22.5)		

## Statistical analyses

<b>Statistical analysis title</b>	Analysis of Overall Survival
Statistical analysis description: A stratified log-rank test was used for the primary comparison of overall survival.	
Comparison groups	Placebo + PLD v Trebananib + PLD
Number of subjects included in analysis	223
Analysis specification	Pre-specified
Analysis type	other <sup>[3]</sup>
P-value	= 0.76 <sup>[4]</sup>
Method	Stratified Log-rank Test

Notes:

[3] - The analysis was descriptive

[4] - Stratified by PFI status and enrollment before PLD shortage.

<b>Statistical analysis title</b>	Cox Proportional Hazards Analysis
Statistical analysis description: A stratified Cox regression model was also used to provide the estimated overall survival hazard ratio and 2-sided 95% CI for trebananib in combination with PLD relative to placebo and PLD.	
Comparison groups	Placebo + PLD v Trebananib + PLD
Number of subjects included in analysis	223
Analysis specification	Pre-specified
Analysis type	other
P-value	= 0.76 <sup>[5]</sup>
Method	Cox proportional hazard model
Parameter estimate	Hazard ratio (HR)
Point estimate	1.04
Confidence interval	
level	95 %
sides	2-sided
lower limit	0.79
upper limit	1.39

Notes:

[5] - Stratified by PFI status and enrollment before PLD shortage

## Secondary: Objective Response Rate

<b>End point title</b>	Objective Response Rate
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End point description:

Disease response was assessed using computed tomography or magnetic resonance imaging of at least the chest, abdomen and pelvis. Objective response rate (ORR) was defined as the percentage of participants with measurable disease at baseline who achieved either a complete response (CR) or partial response (PR) while on study, according to RECIST 1.1 mod assessed by the investigator. CR: Disappearance of all target and non-target lesions and no new lesions. Any pathological lymph nodes must have reduction in short axis to < 10 mm.

PR: At least a 30% decrease in the size of target lesions with persistence of one or more non-target lesions and no new lesions, or, disappearance of all target lesions with persistence of one or more non-target lesions and no new lesions.

Participants with measurable disease at baseline who did not meet the criteria for objective response by the analysis cut-off date were considered non-responders.

End point type	Secondary
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End point timeframe:

Disease response was assessed every 8 weeks for the first 64 weeks, then every 16 weeks for 32 weeks, and every 24 weeks thereafter until disease progression or death.

End point values	Placebo + PLD	Trebananib + PLD		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	94 <sup>[6]</sup>	99 <sup>[7]</sup>		
Units: percentage of participants				
number (confidence interval 95%)	20.2 (12.6 to 29.8)	47.5 (37.3 to 57.8)		

Notes:

[6] - Participants with measurable disease at baseline

[7] - Participants with measurable disease at baseline

## Statistical analyses

Statistical analysis title	Analysis of Objective Response Rate
Comparison groups	Placebo + PLD v Trebananib + PLD
Number of subjects included in analysis	193
Analysis specification	Pre-specified
Analysis type	other
P-value	= 0 <sup>[8]</sup>
Method	Cochran-Mantel-Haenszel
Parameter estimate	Odds ratio (OR)
Point estimate	3.79
Confidence interval	
level	95 %
sides	2-sided
lower limit	1.95
upper limit	7.35

Notes:

[8] - Cochran-Mantel-Haenszel test adjusted for PFI status.

## Secondary: Duration of Response

End point title	Duration of Response
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End point description:

Duration of response was defined as the time from the first objective response to disease progression per RECIST 1.1 with modifications or death due to any cause. Subjects not meeting criteria for disease progression by the analysis data cut-off date were censored at their last evaluable disease assessment date. The analysis of DOR was conducted on the subset of subjects with measurable disease at baseline who experienced an objective response during the study.

End point type	Secondary
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End point timeframe:

Disease response was assessed every 8 weeks for the first 64 weeks, then every 16 weeks for 32 weeks, and every 24 weeks thereafter until disease progression or death.

<b>End point values</b>	Placebo + PLD	Trebananib + PLD		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	19 <sup>[9]</sup>	47 <sup>[10]</sup>		
Units: months				
median (confidence interval 95%)	5.6 (2.3 to 9.2)	7.4 (5.6 to 9.1)		

Notes:

[9] - Participants with measurable disease at baseline and with an objective response during the study

[10] - Participants with measurable disease at baseline and with an objective response during the study

## Statistical analyses

No statistical analyses for this end point

## Secondary: CA-125 Response Rate

End point title	CA-125 Response Rate
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End point description:

A confirmed CA-125 response, according to the Gynecologic Cancer Intergroup (GCIG) criteria, defined as the percentage of participants with at least a 50% reduction in CA-125 levels from baseline, confirmed and maintained for at least 28 days. Only participants with CA-125 levels at least 2 X the upper limit of normal (ULN) within 2 weeks of starting treatment were evaluated for CA-125 response. Participants evaluable for CA-125 response that did not meet the criteria for a CA-125 response were considered non-responders.

End point type	Secondary
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End point timeframe:

CA-125 was measured every 4 weeks for up to 2 years and then every 6 months thereafter.

<b>End point values</b>	Placebo + PLD	Trebananib + PLD		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	87 <sup>[11]</sup>	94 <sup>[12]</sup>		
Units: percentage of participants				
number (confidence interval 95%)	26.4 (17.6 to 37.0)	51.1 (40.5 to 61.5)		

Notes:

[11] - Participants with baseline CA-125 at least 2 x upper limit of normal

[12] - Participants with baseline CA-125 at least 2 x upper limit of normal

## Statistical analyses

<b>Statistical analysis title</b>	Analysis of CA-125 Response
Comparison groups	Placebo + PLD v Trebananib + PLD

Number of subjects included in analysis	181
Analysis specification	Pre-specified
Analysis type	other
P-value	= 0.001 <sup>[13]</sup>
Method	Cochran-Mantel-Haenszel
Parameter estimate	Odds ratio (OR)
Point estimate	3.05
Confidence interval	
level	95 %
sides	2-sided
lower limit	1.61
upper limit	5.79

Notes:

[13] - Cochran-Mantel-Haenszel test adjusted for PFI status.

### Secondary: Maximum Percent Change from Baseline in CA-125

End point title	Maximum Percent Change from Baseline in CA-125
End point description:	
End point type	Secondary
End point timeframe:	
Baseline and every 4 weeks for 2 years and then every 6 months thereafter until the end of treatment.	

End point values	Placebo + PLD	Trebananib + PLD		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	75 <sup>[14]</sup>	85 <sup>[15]</sup>		
Units: percent change				
arithmetic mean (standard error)	-13.22 (± 16.83)	-46.45 (± 9.01)		

Notes:

[14] - Participants with baseline CA-125  $\geq 2 \times$  ULN and available post-baseline data

[15] - Participants with baseline CA-125  $\geq 2 \times$  ULN and available post-baseline data

### Statistical analyses

No statistical analyses for this end point

### Secondary: Number of Participants with Adverse Events

End point title	Number of Participants with Adverse Events
End point description:	
Adverse events were graded for severity using the Common Terminology Criteria for Adverse Events version 3.0.	
Trebananib/placebo-related or PLD-related adverse events are those events for which the investigator considered there to be a reasonable possibility that the event may have been caused by the study treatment, trebananib/placebo or PLD, respectively.	
End point type	Secondary
End point timeframe:	
From the first dose of study drug until 30 days after last dose; the median duration of trebananib treatment was 156 days.	

End point values	Placebo + PLD	Trebananib + PLD		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	108 <sup>[16]</sup>	113 <sup>[17]</sup>		
Units: participants				
Any adverse event	108	113		
AE Grade $\geq 3$	78	89		
AE Grade $\geq 4$	21	15		
Fatal adverse events	7	6		
Serious adverse events	50	55		
AE leading to discontinuation of trebananib	23	32		
AE leading to discontinuation of PLD	25	20		
AE leading to discontinuation from study treatment	16	12		
Trebananib/placebo-related adverse events	89	95		
Trebananib/placebo-related AE Grade $\geq 3$	33	48		
Trebananib/placebo-related AE Grade $\geq 4$	8	6		
Trebananib/placebo-related fatal AE	2	2		
Trebananib/placebo-related serious AE	18	27		
PLD-related adverse events	99	105		
PLD-related adverse events Grade $\geq 3$	44	53		
PLD-related adverse events Grade $\geq 4$	8	6		
PLD-related fatal adverse events	0	0		
PLD-related serious adverse events	14	17		

Notes:

[16] - Randomized participants who received at least 1 dose of trebananib or PLD

[17] - Randomized participants who received at least 1 dose of trebananib or PLD

## Statistical analyses

No statistical analyses for this end point

## Secondary: Maximum Observed Serum Concentration (Cmax) of Trebananib and Week 1 and Week 5

End point title	Maximum Observed Serum Concentration (Cmax) of Trebananib and Week 1 and Week 5
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End point description:

End point type	Secondary
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End point timeframe:

Week 1 and week 5 at the end of infusion

End point values	Placebo + PLD	Trebananib + PLD		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	0 <sup>[18]</sup>	101 <sup>[19]</sup>		
Units: µg/mL				
arithmetic mean (standard deviation)				
Week 1	( )	295 (± 215)		
Week 5	( )	319 (± 105)		

Notes:

[18] - Participants did not receive trebananib

[19] - Participants with evaluable trebananib concentration data; N= 70 at Week 5

### Statistical analyses

No statistical analyses for this end point

### Secondary: Minimum Observed Serum Concentration (Cmin) of Trebananib and Weeks 2, 5 and 9

End point title	Minimum Observed Serum Concentration (Cmin) of Trebananib and Weeks 2, 5 and 9
End point description:	
End point type	Secondary
End point timeframe:	
Weeks 2, 5 and 9, predose	

End point values	Placebo + PLD	Trebananib + PLD		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	0 <sup>[20]</sup>	101 <sup>[21]</sup>		
Units: µg/mL				
arithmetic mean (standard deviation)				
Week 2	( )	13.2 (± 6.56)		
Week 5	( )	28.2 (± 15.8)		
Week 9	( )	27.8 (± 15.5)		

Notes:

[20] - Participants did not receive trebananib

[21] - Participants with evaluable data; N at weeks 5 and 9 was 76 and 67

### Statistical analyses

No statistical analyses for this end point

### Secondary: Number of Participants who Developed Anti-trebananib Antibodies

End point title	Number of Participants who Developed Anti-trebananib Antibodies
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End point description:

Two validated assays were used to detect the presence of anti-trebananib antibodies. Samples were first tested in a biosensor immunoassay to detect antibodies capable of binding to trebananib. Samples confirmed to be positive for binding antibodies were subsequently tested in a receptor-binding assay to determine neutralizing activity against trebananib. If a post-dose sample was positive for binding antibodies and demonstrated neutralizing activity at the same time point, the sample was defined as

positive for neutralizing antibodies.

End point type	Secondary
End point timeframe:	
Pre-infusion of trebananib or placebo on day 1 of week 1, week 9 and at the safety follow-up visit.	

End point values	Placebo + PLD	Trebananib + PLD		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	95 <sup>[22]</sup>	101 <sup>[23]</sup>		
Units: participants				
Binding antibody positive	0	1		
Neutralizing antibody positive	0	0		

Notes:

[22] - Participants with a postbaseline result

[23] - Participants with a postbaseline result

## Statistical analyses

No statistical analyses for this end point

## Secondary: Secondary: Change from Baseline in Functional Assessment of Cancer Therapy – Ovary (FACT-O) Summary Score

End point title	Secondary: Change from Baseline in Functional Assessment of Cancer Therapy – Ovary (FACT-O) Summary Score
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End point description:

The FACT-O evaluates the health-related quality of life (HRQOL) and symptoms in patients with ovarian cancer. It consists of the FACT-G, a 27 item general cancer questionnaire and a 12-item ovarian cancer-specific subscale (OCS). Each item is scored by the participant on a scale from 0 (not at all true) to 4 (very much true). The total score ranges from 0 to 156; a higher total score indicates better quality of life or less severe symptoms.

The patient-reported outcomes (PRO) analysis set is defined as a subset of randomized subjects who have a baseline PRO assessment and at least one post-baseline PRO assessment prior to disease progression.

End point type	Secondary
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End point timeframe:

Baseline and Weeks 5, 9, 13, 17, 25, 33, 41, 49, 57, 65, 81, and the safety follow-up visit 30 days after last dose.

End point values	Placebo + PLD	Trebananib + PLD		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	90 <sup>[24]</sup>	98 <sup>[25]</sup>		
Units: units on a scale				
arithmetic mean (standard deviation)				
Week 5 (n = 87, 91)	-2.495 (± 19.732)	-4.991 (± 13.523)		
Week 9 (n = 69, 82)	-0.116 (± 17.978)	-5.708 (± 14.545)		
Week 13 (n = 62, 72)	-2.502 (± 18.251)	-7.161 (± 17.231)		



Week 17 (n = 52, 61)	-1.554 (± 16.545)	-5.809 (± 18.753)		
Week 25 (n = 42, 44)	1.568 (± 15.889)	-5.093 (± 15.862)		
Week 33 (n = 30, 29)	0.926 (± 14.907)	0.204 (± 11.304)		
Week 41 (n = 21, 21)	1.622 (± 14.203)	-0.192 (± 1.474)		
Week 49 (n = 12, 11)	2.113 (± 12.096)	-3.970 (± 15.934)		
Week 57 (n = 9, 4)	9.333 (± 13.043)	-6.733 (± 13.244)		
Week 65 (n = 8, 4)	3.621 (± 12.060)	-2.833 (± 12.662)		
Week 81 (n = 5, 2)	11.267 (± 7.316)	1.500 (± 2.121)		
Safety follow-up (n = 13, 14)	-11.667 (± 21.698)	-8.446 (± 18.886)		

Notes:

[24] - Participants with available baseline and post-baseline FACT-O data

[25] - Participants with available baseline and post-baseline FACT-O data

## Statistical analyses

No statistical analyses for this end point

## Secondary: Change from Baseline in FACT-O Ovarian Cancer-specific (OCS) Subscale

End point title	Change from Baseline in FACT-O Ovarian Cancer-specific (OCS) Subscale
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End point description:

The FACT-O evaluates the health-related quality of life (HRQOL) and symptoms in patients with ovarian cancer. It consists of the FACT-G, a 27 item general cancer questionnaire and a 12-item ovarian cancer-specific subscale (OCS). The OCS consists of 12 symptom items including swelling in stomach area, cramps in stomach area, weight loss, hair loss, control of bowels, appetite, vomiting, ability to get around, liking the appearance of one's body, being able to feel like a woman, interest in sex, and concern about ability to have children. Each item is scored by the participant on a scale from 0 (not at all true) to 4 (very much true). The OCS summary score ranges from 0 to 48, where a higher score indicates better quality of life or less severe symptoms.

The PRO analysis set includes a subset of randomized subjects who had a baseline PRO assessment and at least one post-baseline PRO assessment prior to disease progression.

End point type	Secondary
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End point timeframe:

Baseline and Weeks 5, 9, 13, 17, 25, 33, 41, 49, 57, 65, 81, and the safety follow-up visit 30 days after last dose.

End point values	Placebo + PLD	Trebananib + PLD		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	91 <sup>[26]</sup>	98 <sup>[27]</sup>		
Units: units on a scale				
arithmetic mean (standard deviation)				
Week 5 (n = 89, 91)	0.129 (± 6.737)	-1.430 (± 4.746)		
Week 9 (n = 71, 83)	0.494 (± 6.173)	-0.993 (± 4.999)		

Week 13 (n = 62, 73)	0.681 (± 6.341)	-1.377 (± 5.325)		
Week 17 (n = 53, 62)	0.752 (± 6.062)	-1.030 (± 6.915)		
Week 25 (n = 42, 45)	0.637 (± 6.187)	-1.108 (± 4.585)		
Week 33 (n = 30, 29)	1.207 (± 5.599)	0.497 (± 4.538)		
Week 41 (n = 21, 22)	2.917 (± 5.465)	0.415 (± 4.569)		
Week 49 (n = 12, 11)	1.296 (± 5.465)	-0.182 (± 5.528)		
Week 57 (n = 9, 4)	4.778 (± 6.741)	-1.525 (± 1.684)		
Week 65 (n = 8, 4)	4.875 (± 8.288)	-0.500 (± 2.380)		
Week 81 (n = 5, 2)	7.200 (± 7.918)	-1.500 (± 0.707)		
Safety follow-up (n = 14, 14)	-4.021 (± 8.205)	-3.250 (± 7.350)		

Notes:

[26] - Participants with available baseline and post-baseline FACT-OCS data

[27] - Participants with available baseline and post-baseline FACT-OCS data

## Statistical analyses

No statistical analyses for this end point

## Secondary: Change from Baseline in EuroQOL 5-Dimension (EQ-5D) Health Index Score

End point title	Change from Baseline in EuroQOL 5-Dimension (EQ-5D) Health Index Score
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End point description:

The EQ-5D is a standardized instrument for use as a generic, preference-based measure of health outcome. The EQ-5D questionnaire captures two basic types of information, a descriptive "profile," or "health state," and an overall health rating using a visual analogue scale.

The health state includes five dimensions: mobility, self-care, usual activities, pain/discomfort, and anxiety/depression. Each question has 3 answer choices: 1 (no problems), 2 (moderate problems), and 3 (extreme problems). The health states for each respondent are converted into a single index number using a specified set of weights. Resulting scores can range from 1.0 and -0.594. A higher score indicates a more preferred health status with 1.0 representing perfect health.

The PRO analysis set includes a subset of randomized subjects who had a baseline assessment and at least 1 post-baseline PRO assessment prior to disease progression.

End point type	Secondary
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End point timeframe:

Baseline and Weeks 5, 9, 13, 17, 25, 33, 41, 49, 57, 65, 81, and the safety follow-up visit 30 days after last dose.

End point values	Placebo + PLD	Trebananib + PLD		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	86 <sup>[28]</sup>	93 <sup>[29]</sup>		
Units: units on a scale				
arithmetic mean (standard deviation)				
Week 5 (n = 82, 83)	-0.027 (± 0.217)	-0.026 (± 0.198)		

Week 9 (n = 66, 77)	-0.057 (± 0.237)	-0.035 (± 0.263)		
Week 13 (n = 55, 69)	-0.001 (± 0.210)	-0.044 (± 0.225)		
Week 17 (n = 45, 58)	-0.043 (± 0.178)	-0.052 (± 0.269)		
Week 25 (n = 38, 39)	-0.017 (± 0.155)	-0.050 (± 0.266)		
Week 33 (n = 26, 26)	-0.015 (± 0.188)	-0.061 (± 0.206)		
Week 41 (n = 18, 20)	-0.007 (± 0.177)	-0.078 (± 0.188)		
Week 49 (n = 10, 9)	0.008 (± 0.155)	-0.166 (± 0.319)		
Week 57 (n = 8, 4)	0.060 (± 0.164)	-0.010 (± 0.020)		
Week 65 (n = 7, 4)	0.059 (± 0.178)	-0.010 (± 0.020)		
Week 81 (n = 4, 2)	0.065 (± 0.241)	0.000 (± 0.000)		
Safety follow-up (n = 13, 14)	-0.098 (± 0.158)	-0.159 (± 0.339)		

Notes:

[28] - Participants with available baseline and post-baseline EQ-5D data

[29] - Participants with available baseline and post-baseline EQ-5D data

## Statistical analyses

No statistical analyses for this end point

## Secondary: Change from Baseline in EQ-5D Visual Analogue Scale Score

End point title	Change from Baseline in EQ-5D Visual Analogue Scale Score
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End point description:

The EQ-5D is a standardized instrument for use as a generic, preference-based measure of health outcome. The EQ-5D questionnaire captures two basic types of information, a descriptive "profile," or "health state," and an overall health rating using a visual analogue scale (VAS).

The visual analogue scale asks respondents to rate their present health status on a 0 - 100 visual analogue scale, with 0 labeled as "Worst imaginable health state" and 100 labeled as "Best imaginable health state." The VAS score is determined by observing the point at which the subjects hand drawn line intersects the scale.

The PRO analysis set includes the subset of randomized subjects who had a baseline assessment and at least 1 post-baseline PRO assessment prior to disease progression.

End point type	Secondary
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End point timeframe:

Baseline and Weeks 5, 9, 13, 17, 25, 33, 41, 49, 57, 65, 81, and the safety follow-up visit 30 days after last dose.

End point values	Placebo + PLD	Trebananib + PLD		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	90 <sup>[30]</sup>	94 <sup>[31]</sup>		
Units: units on a scale				
arithmetic mean (standard deviation)				
Week 5 (n = 86, 82)	-3.907 (± 21.469)	-0.622 (± 18.844)		

Week 9 (n = 69, 79)	-3.275 (± 15.224)	0.532 (± 19.367)		
Week 13 (n = 60, 69)	-2.433 (± 16.213)	-0.942 (± 17.306)		
Week 17 (n = 51, 59)	-1.569 (± 15.263)	-1.441 (± 23.219)		
Week 25 (n = 40, 44)	1.125 (± 10.013)	-0.773 (± 21.431)		
Week 33 (n = 28, 28)	2.607 (± 12.467)	4.714 (± 15.224)		
Week 41 (n = 21, 22)	2.000 (± 12.542)	5.682 (± 18.198)		
Week 49 (n = 12, 11)	-2.083 (± 14.656)	1.909 (± 13.472)		
Week 57 (n = 9, 4)	5.000 (± 10.770)	-4.000 (± 4.899)		
Week 65 (n = 8, 4)	1.625 (± 12.727)	-0.750 (± 1.500)		
Week 81 (n = 5, 2)	-0.200 (± 10.257)	0.000 (± 0.000)		
Safety follow-up (n = 15, 14)	-5.867 (± 11.154)	-7.571 (± 18.029)		

Notes:

[30] - Participants with available baseline and post-baseline EQ-5D VAS data

[31] - Participants with available baseline and post-baseline EQ-5D VAS data

## Statistical analyses

No statistical analyses for this end point

## Adverse events

### Adverse events information

Timeframe for reporting adverse events:

From the first dose of study drug until 30 days after last dose; the median duration of trebananib treatment was 156 days.

Adverse event reporting additional description:

One subject randomized to the placebo group received trebananib and was counted in the trebananib group for safety analyses.

Assessment type	Systematic
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### Dictionary used

Dictionary name	MedDRA
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Dictionary version	20.0
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### Reporting groups

Reporting group title	Trebananib + PLD
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Reporting group description:

Participants received PLD 50 mg/m<sup>2</sup> every 4 weeks (Q4W) plus trebananib 15 mg/kg administered intravenously (IV) every week (QW) until disease progression per RECIST 1.1 mod, unacceptable toxicity, withdrawal of consent, or death.

Reporting group title	Placebo + PLD
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Reporting group description:

Participants received pegylated liposomal doxorubicin (PLD) 50 mg/m<sup>2</sup> every 4 weeks (Q4W) plus placebo to trebananib administered intravenously (IV) every week (QW) until disease progression per RECIST 1.1 mod, unacceptable toxicity, withdrawal of consent, or death.

Serious adverse events	Trebananib + PLD	Placebo + PLD	
Total subjects affected by serious adverse events			
subjects affected / exposed	55 / 113 (48.67%)	50 / 108 (46.30%)	
number of deaths (all causes)	102	93	
number of deaths resulting from adverse events			
Neoplasms benign, malignant and unspecified (incl cysts and polyps)			
Adenocarcinoma			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 1	0 / 0	
Malignant ascites			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Malignant pleural effusion			

subjects affected / exposed	1 / 113 (0.88%)	2 / 108 (1.85%)	
occurrences causally related to treatment / all	0 / 1	1 / 3	
deaths causally related to treatment / all	0 / 0	0 / 0	
Metastases to central nervous system			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	0 / 2	0 / 0	
deaths causally related to treatment / all	0 / 1	0 / 0	
Metastases to meninges			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Ovarian cancer metastatic			
subjects affected / exposed	0 / 113 (0.00%)	3 / 108 (2.78%)	
occurrences causally related to treatment / all	0 / 0	0 / 3	
deaths causally related to treatment / all	0 / 0	0 / 2	
Vascular disorders			
Deep vein thrombosis			
subjects affected / exposed	2 / 113 (1.77%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	1 / 2	1 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Embolism			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hypotension			
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	1 / 1	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Peripheral venous disease			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Venous thrombosis limb			

subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	2 / 2	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
General disorders and administration site conditions			
Asthenia			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Chest discomfort			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Chest pain			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Chills			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Complication associated with device			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
General physical health deterioration			
subjects affected / exposed	1 / 113 (0.88%)	2 / 108 (1.85%)	
occurrences causally related to treatment / all	0 / 2	0 / 2	
deaths causally related to treatment / all	0 / 0	0 / 0	
Generalised oedema			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Localised oedema			

subjects affected / exposed	2 / 113 (1.77%)	2 / 108 (1.85%)	
occurrences causally related to treatment / all	1 / 2	3 / 3	
deaths causally related to treatment / all	0 / 0	0 / 0	
Mucosal inflammation			
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 1	1 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Multiple organ dysfunction syndrome			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 1	
Pain			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pyrexia			
subjects affected / exposed	5 / 113 (4.42%)	4 / 108 (3.70%)	
occurrences causally related to treatment / all	0 / 6	0 / 4	
deaths causally related to treatment / all	0 / 0	0 / 0	
Suprapubic pain			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Immune system disorders			
Anaphylactic reaction			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hypersensitivity			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Respiratory, thoracic and mediastinal disorders			



Alveolitis allergic			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Bronchospasm			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	1 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Cough			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Dyspnoea			
subjects affected / exposed	3 / 113 (2.65%)	2 / 108 (1.85%)	
occurrences causally related to treatment / all	1 / 3	2 / 4	
deaths causally related to treatment / all	0 / 0	0 / 1	
Hypoxia			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	2 / 2	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pleural effusion			
subjects affected / exposed	8 / 113 (7.08%)	4 / 108 (3.70%)	
occurrences causally related to treatment / all	6 / 8	2 / 6	
deaths causally related to treatment / all	0 / 0	0 / 1	
Pleurisy			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pneumonia aspiration			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pneumothorax			

subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	1 / 1	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pulmonary embolism			
subjects affected / exposed	2 / 113 (1.77%)	3 / 108 (2.78%)	
occurrences causally related to treatment / all	1 / 2	4 / 4	
deaths causally related to treatment / all	0 / 0	1 / 1	
Respiratory failure			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	1 / 1	
deaths causally related to treatment / all	0 / 0	1 / 1	
Psychiatric disorders			
Confusional state			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Product issues			
Device malfunction			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Investigations			
Blood creatine increased			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Blood creatinine abnormal			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Eastern cooperative oncology group performance status worsened			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	

Weight decreased subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Injury, poisoning and procedural complications			
Gastrointestinal stoma complication subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	1 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Infusion related reaction subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Cardiac disorders			
Cardiac failure subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pericardial effusion subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Right ventricular failure subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	1 / 1	0 / 0	
Sinus tachycardia subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Tachycardia subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	

Nervous system disorders			
Cerebral ischaemia			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	2 / 2	0 / 0	
deaths causally related to treatment / all	1 / 1	0 / 0	
Blood and lymphatic system disorders			
Anaemia			
subjects affected / exposed	2 / 113 (1.77%)	2 / 108 (1.85%)	
occurrences causally related to treatment / all	0 / 6	0 / 3	
deaths causally related to treatment / all	0 / 0	0 / 0	
Febrile neutropenia			
subjects affected / exposed	0 / 113 (0.00%)	3 / 108 (2.78%)	
occurrences causally related to treatment / all	0 / 0	2 / 3	
deaths causally related to treatment / all	0 / 0	0 / 0	
Thrombocytopenia			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Gastrointestinal disorders			
Abdominal distension			
subjects affected / exposed	0 / 113 (0.00%)	3 / 108 (2.78%)	
occurrences causally related to treatment / all	0 / 0	2 / 3	
deaths causally related to treatment / all	0 / 0	0 / 0	
Abdominal pain			
subjects affected / exposed	7 / 113 (6.19%)	6 / 108 (5.56%)	
occurrences causally related to treatment / all	2 / 11	1 / 6	
deaths causally related to treatment / all	0 / 0	0 / 0	
Abdominal pain upper			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Ascites			

subjects affected / exposed	11 / 113 (9.73%)	5 / 108 (4.63%)	
occurrences causally related to treatment / all	14 / 23	3 / 5	
deaths causally related to treatment / all	0 / 0	0 / 0	
Constipation			
subjects affected / exposed	1 / 113 (0.88%)	2 / 108 (1.85%)	
occurrences causally related to treatment / all	0 / 2	0 / 3	
deaths causally related to treatment / all	0 / 0	0 / 0	
Diarrhoea			
subjects affected / exposed	3 / 113 (2.65%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 5	1 / 1	
deaths causally related to treatment / all	0 / 1	0 / 0	
Enteritis			
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	0 / 2	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Erosive oesophagitis			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	1 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Faecaloma			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Gastrointestinal oedema			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	2 / 2	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Gastrointestinal perforation			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Gastrooesophageal reflux disease			

subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Gingival pain			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Ileus			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Intestinal obstruction			
subjects affected / exposed	2 / 113 (1.77%)	6 / 108 (5.56%)	
occurrences causally related to treatment / all	0 / 2	0 / 7	
deaths causally related to treatment / all	0 / 0	0 / 0	
Intra-abdominal fluid collection			
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	3 / 4	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Large intestinal obstruction			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Nausea			
subjects affected / exposed	3 / 113 (2.65%)	4 / 108 (3.70%)	
occurrences causally related to treatment / all	3 / 4	1 / 5	
deaths causally related to treatment / all	0 / 0	0 / 0	
Odynophagia			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	1 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Oesophagitis			

subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Rectal haemorrhage			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Small intestinal obstruction			
subjects affected / exposed	4 / 113 (3.54%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 4	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Stomatitis			
subjects affected / exposed	3 / 113 (2.65%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	1 / 3	1 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Stomatitis necrotising			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Subileus			
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	1 / 1	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Vomiting			
subjects affected / exposed	9 / 113 (7.96%)	3 / 108 (2.78%)	
occurrences causally related to treatment / all	4 / 14	0 / 4	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hepatobiliary disorders			
Portal vein thrombosis			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	1 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Skin and subcutaneous tissue disorders			

Dermatitis allergic			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Palmar-plantar erythrodysaesthesia syndrome			
subjects affected / exposed	1 / 113 (0.88%)	2 / 108 (1.85%)	
occurrences causally related to treatment / all	1 / 1	1 / 2	
deaths causally related to treatment / all	0 / 0	0 / 0	
Urticaria			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Renal and urinary disorders			
Hydronephrosis			
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	0 / 2	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Urinary tract obstruction			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Musculoskeletal and connective tissue disorders			
Arthralgia			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pain in extremity			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Infections and infestations			
Bacteraemia			



subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Bronchitis			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Catheter site infection			
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 1	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Device related infection			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Influenza			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Neutropenic sepsis			
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 1	1 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pneumonia			
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 1	1 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pyelonephritis			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences causally related to treatment / all	0 / 0	0 / 2	
deaths causally related to treatment / all	0 / 0	0 / 0	
Urinary tract infection enterococcal			

subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Viral infection			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Metabolism and nutrition disorders			
Cachexia			
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	0 / 2	0 / 0	
deaths causally related to treatment / all	0 / 1	0 / 0	
Decreased appetite			
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	1 / 2	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Dehydration			
subjects affected / exposed	1 / 113 (0.88%)	2 / 108 (1.85%)	
occurrences causally related to treatment / all	1 / 1	0 / 2	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hyperglycaemia			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hypokalaemia			
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	2 / 2	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hypovolaemia			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	

Frequency threshold for reporting non-serious adverse events: 0 %

<b>Non-serious adverse events</b>	Trebananib + PLD	Placebo + PLD	
Total subjects affected by non-serious adverse events			
subjects affected / exposed	112 / 113 (99.12%)	103 / 108 (95.37%)	
Neoplasms benign, malignant and unspecified (incl cysts and polyps)			
Malignant pleural effusion			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Pyogenic granuloma			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Tumour pain			
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)	
occurrences (all)	1	1	
Vascular disorders			
Aortic thrombosis			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Deep vein thrombosis			
subjects affected / exposed	4 / 113 (3.54%)	1 / 108 (0.93%)	
occurrences (all)	4	1	
Diastolic hypertension			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Flushing			
subjects affected / exposed	3 / 113 (2.65%)	4 / 108 (3.70%)	
occurrences (all)	3	4	
Embolism			
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)	
occurrences (all)	1	1	
Haematoma			
subjects affected / exposed	2 / 113 (1.77%)	2 / 108 (1.85%)	
occurrences (all)	2	3	
Hot flush			

subjects affected / exposed	5 / 113 (4.42%)	1 / 108 (0.93%)	
occurrences (all)	5	1	
Hypertension			
subjects affected / exposed	12 / 113 (10.62%)	9 / 108 (8.33%)	
occurrences (all)	17	9	
Jugular vein distension			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Hypotension			
subjects affected / exposed	0 / 113 (0.00%)	2 / 108 (1.85%)	
occurrences (all)	0	2	
Jugular vein thrombosis			
subjects affected / exposed	2 / 113 (1.77%)	1 / 108 (0.93%)	
occurrences (all)	2	1	
Lymphocele			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Peripheral venous disease			
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)	
occurrences (all)	1	1	
Lymphoedema			
subjects affected / exposed	10 / 113 (8.85%)	2 / 108 (1.85%)	
occurrences (all)	16	2	
Phlebitis			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Thrombosis			
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)	
occurrences (all)	2	0	
Thrombophlebitis			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Venous thrombosis			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Surgical and medical procedures			

Central venous catheter removal subjects affected / exposed occurrences (all)	1 / 113 (0.88%) 1	0 / 108 (0.00%) 0	
Tooth extraction subjects affected / exposed occurrences (all)	1 / 113 (0.88%) 1	0 / 108 (0.00%) 0	
Eye irrigation subjects affected / exposed occurrences (all)	1 / 113 (0.88%) 1	0 / 108 (0.00%) 0	
General disorders and administration site conditions			
Application site hypersensitivity subjects affected / exposed occurrences (all)	0 / 113 (0.00%) 0	1 / 108 (0.93%) 1	
Asthenia subjects affected / exposed occurrences (all)	9 / 113 (7.96%) 16	12 / 108 (11.11%) 19	
Axillary pain subjects affected / exposed occurrences (all)	2 / 113 (1.77%) 2	0 / 108 (0.00%) 0	
Catheter site erythema subjects affected / exposed occurrences (all)	1 / 113 (0.88%) 2	1 / 108 (0.93%) 4	
Catheter site haematoma subjects affected / exposed occurrences (all)	1 / 113 (0.88%) 1	0 / 108 (0.00%) 0	
Catheter site inflammation subjects affected / exposed occurrences (all)	0 / 113 (0.00%) 0	1 / 108 (0.93%) 1	
Catheter site pain subjects affected / exposed occurrences (all)	2 / 113 (1.77%) 3	1 / 108 (0.93%) 1	
Catheter site pruritus subjects affected / exposed occurrences (all)	0 / 113 (0.00%) 0	1 / 108 (0.93%) 3	
Catheter site rash			

subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Catheter site vesicles		
subjects affected / exposed	1 / 113 (0.88%)	2 / 108 (1.85%)
occurrences (all)	1	2
Chest discomfort		
subjects affected / exposed	2 / 113 (1.77%)	1 / 108 (0.93%)
occurrences (all)	2	1
Chest pain		
subjects affected / exposed	2 / 113 (1.77%)	5 / 108 (4.63%)
occurrences (all)	2	5
Chills		
subjects affected / exposed	7 / 113 (6.19%)	2 / 108 (1.85%)
occurrences (all)	10	5
Complication associated with device		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Discomfort		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Drug intolerance		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Early satiety		
subjects affected / exposed	0 / 113 (0.00%)	4 / 108 (3.70%)
occurrences (all)	0	5
Fatigue		
subjects affected / exposed	61 / 113 (53.98%)	48 / 108 (44.44%)
occurrences (all)	112	105
Gait disturbance		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	1
General physical health deterioration		
subjects affected / exposed	3 / 113 (2.65%)	1 / 108 (0.93%)
occurrences (all)	3	1
Generalised oedema		

subjects affected / exposed	8 / 113 (7.08%)	2 / 108 (1.85%)
occurrences (all)	13	2
Hernia pain		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	1
Impaired healing		
subjects affected / exposed	1 / 113 (0.88%)	2 / 108 (1.85%)
occurrences (all)	1	11
Influenza like illness		
subjects affected / exposed	6 / 113 (5.31%)	2 / 108 (1.85%)
occurrences (all)	7	2
Infusion site extravasation		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Injection site haematoma		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Injection site rash		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Injection site reaction		
subjects affected / exposed	1 / 113 (0.88%)	2 / 108 (1.85%)
occurrences (all)	1	2
Localised oedema		
subjects affected / exposed	67 / 113 (59.29%)	32 / 108 (29.63%)
occurrences (all)	133	45
Malaise		
subjects affected / exposed	5 / 113 (4.42%)	2 / 108 (1.85%)
occurrences (all)	5	2
Mucosal dryness		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Mucosal inflammation		
subjects affected / exposed	20 / 113 (17.70%)	26 / 108 (24.07%)
occurrences (all)	40	64
Non-cardiac chest pain		

subjects affected / exposed	4 / 113 (3.54%)	1 / 108 (0.93%)	
occurrences (all)	5	1	
Oedema peripheral			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	3	0	
Pain			
subjects affected / exposed	4 / 113 (3.54%)	8 / 108 (7.41%)	
occurrences (all)	4	12	
Peripheral swelling			
subjects affected / exposed	1 / 113 (0.88%)	2 / 108 (1.85%)	
occurrences (all)	1	2	
Pyrexia			
subjects affected / exposed	20 / 113 (17.70%)	15 / 108 (13.89%)	
occurrences (all)	28	19	
Sensation of foreign body			
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)	
occurrences (all)	1	1	
Systemic inflammatory response syndrome			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Temperature intolerance			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Ulcer			
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)	
occurrences (all)	2	0	
Visceral oedema			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Immune system disorders			
Anaphylactic reaction			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Drug hypersensitivity			



subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)	
occurrences (all)	1	1	
Hypersensitivity			
subjects affected / exposed	2 / 113 (1.77%)	5 / 108 (4.63%)	
occurrences (all)	2	5	
Seasonal allergy			
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)	
occurrences (all)	3	0	
Reproductive system and breast disorders			
Genital rash			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Pelvic discomfort			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Pelvic pain			
subjects affected / exposed	2 / 113 (1.77%)	1 / 108 (0.93%)	
occurrences (all)	2	1	
Vaginal discharge			
subjects affected / exposed	3 / 113 (2.65%)	1 / 108 (0.93%)	
occurrences (all)	4	2	
Vaginal haemorrhage			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	2	
Vaginal lesion			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	2	
Vulva cyst			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Vulval disorder			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	2	0	
Vulvovaginal discomfort			

subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Vulvovaginal dryness			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Vulvovaginal inflammation			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Vulvovaginal pruritus			
subjects affected / exposed	3 / 113 (2.65%)	1 / 108 (0.93%)	
occurrences (all)	3	1	
Respiratory, thoracic and mediastinal disorders			
Cough			
subjects affected / exposed	22 / 113 (19.47%)	17 / 108 (15.74%)	
occurrences (all)	30	25	
Dysphonia			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Dyspnoea			
subjects affected / exposed	25 / 113 (22.12%)	16 / 108 (14.81%)	
occurrences (all)	51	23	
Dyspnoea at rest			
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)	
occurrences (all)	1	1	
Dyspnoea exertional			
subjects affected / exposed	3 / 113 (2.65%)	6 / 108 (5.56%)	
occurrences (all)	4	9	
Epistaxis			
subjects affected / exposed	3 / 113 (2.65%)	3 / 108 (2.78%)	
occurrences (all)	3	4	
Haemoptysis			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Hiccups			

subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	1
Hypoxia		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Nasal congestion		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	1
Nasal dryness		
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)
occurrences (all)	2	0
Oropharyngeal pain		
subjects affected / exposed	15 / 113 (13.27%)	11 / 108 (10.19%)
occurrences (all)	16	13
Pleural effusion		
subjects affected / exposed	13 / 113 (11.50%)	7 / 108 (6.48%)
occurrences (all)	14	8
Pneumonitis		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	1
Pneumothorax		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Pulmonary artery thrombosis		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Productive cough		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Pulmonary embolism		
subjects affected / exposed	1 / 113 (0.88%)	2 / 108 (1.85%)
occurrences (all)	1	2
Pulmonary pain		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Pulmonary sarcoidosis		

subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Respiratory alkalosis			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Rhinitis allergic			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Rhinorrhoea			
subjects affected / exposed	5 / 113 (4.42%)	1 / 108 (0.93%)	
occurrences (all)	5	1	
Sinus congestion			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Throat irritation			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Psychiatric disorders			
Anxiety			
subjects affected / exposed	7 / 113 (6.19%)	5 / 108 (4.63%)	
occurrences (all)	7	7	
Anxiety disorder			
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)	
occurrences (all)	2	0	
Confusional state			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	2	
Depressed mood			
subjects affected / exposed	0 / 113 (0.00%)	2 / 108 (1.85%)	
occurrences (all)	0	2	
Depression			
subjects affected / exposed	10 / 113 (8.85%)	3 / 108 (2.78%)	
occurrences (all)	10	3	
Disorientation			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	

Hallucination			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Insomnia			
subjects affected / exposed	13 / 113 (11.50%)	11 / 108 (10.19%)	
occurrences (all)	14	14	
Mood altered			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Mood swings			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Panic attack			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Restlessness			
subjects affected / exposed	0 / 113 (0.00%)	2 / 108 (1.85%)	
occurrences (all)	0	2	
Somnambulism			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Stress			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Product issues			
Device dislocation			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Thrombosis in device			
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)	
occurrences (all)	1	2	
Investigations			
Alanine aminotransferase increased			
subjects affected / exposed	3 / 113 (2.65%)	3 / 108 (2.78%)	
occurrences (all)	3	4	
Aspartate aminotransferase increased			

subjects affected / exposed	2 / 113 (1.77%)	4 / 108 (3.70%)
occurrences (all)	2	6
Blood albumin decreased		
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)
occurrences (all)	3	0
Blood alkaline phosphatase increased		
subjects affected / exposed	3 / 113 (2.65%)	1 / 108 (0.93%)
occurrences (all)	5	1
Blood creatinine increased		
subjects affected / exposed	3 / 113 (2.65%)	1 / 108 (0.93%)
occurrences (all)	3	1
Blood folate decreased		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Blood magnesium decreased		
subjects affected / exposed	2 / 113 (1.77%)	2 / 108 (1.85%)
occurrences (all)	2	2
Blood potassium decreased		
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)
occurrences (all)	2	0
Blood potassium increased		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Blood pressure increased		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
C-reactive protein increased		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	1
Eastern cooperative oncology group performance status		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Ejection fraction decreased		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0

Electrocardiogram qt prolonged		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Gamma-glutamyltransferase increased		
subjects affected / exposed	1 / 113 (0.88%)	2 / 108 (1.85%)
occurrences (all)	1	2
Glomerular filtration rate decreased		
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)
occurrences (all)	2	0
Haemoglobin decreased		
subjects affected / exposed	3 / 113 (2.65%)	2 / 108 (1.85%)
occurrences (all)	4	2
Human epidermal growth factor receptor decreased		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Intraocular pressure increased		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	2	0
Lymphocyte count decreased		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	15	1
Neutrophil count decreased		
subjects affected / exposed	5 / 113 (4.42%)	6 / 108 (5.56%)
occurrences (all)	36	7
Platelet count abnormal		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Platelet count decreased		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	5	3
Platelet count increased		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Weight decreased		

subjects affected / exposed occurrences (all)	4 / 113 (3.54%) 4	11 / 108 (10.19%) 17	
Weight increased subjects affected / exposed occurrences (all)	2 / 113 (1.77%) 3	0 / 108 (0.00%) 0	
White blood cell count decreased subjects affected / exposed occurrences (all)	3 / 113 (2.65%) 33	2 / 108 (1.85%) 4	
Injury, poisoning and procedural complications			
Arthropod bite subjects affected / exposed occurrences (all)	1 / 113 (0.88%) 1	0 / 108 (0.00%) 0	
Contusion subjects affected / exposed occurrences (all)	2 / 113 (1.77%) 2	3 / 108 (2.78%) 4	
Eye injury subjects affected / exposed occurrences (all)	0 / 113 (0.00%) 0	1 / 108 (0.93%) 1	
Fall subjects affected / exposed occurrences (all)	0 / 113 (0.00%) 0	3 / 108 (2.78%) 3	
Humerus fracture subjects affected / exposed occurrences (all)	0 / 113 (0.00%) 0	1 / 108 (0.93%) 1	
Infusion related reaction subjects affected / exposed occurrences (all)	0 / 113 (0.00%) 0	3 / 108 (2.78%) 3	
Joint dislocation subjects affected / exposed occurrences (all)	1 / 113 (0.88%) 1	0 / 108 (0.00%) 0	
Laceration subjects affected / exposed occurrences (all)	1 / 113 (0.88%) 1	0 / 108 (0.00%) 0	
Ligament sprain			



subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)
occurrences (all)	2	0
Procedural dizziness		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Procedural pain		
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)
occurrences (all)	3	0
Procedural pneumothorax		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Radiation skin injury		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Rib fracture		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Skin abrasion		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	1
Stoma site discomfort		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Stoma site irritation		
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)
occurrences (all)	3	0
Tendon rupture		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Thermal burn		
subjects affected / exposed	1 / 113 (0.88%)	2 / 108 (1.85%)
occurrences (all)	3	2
Vaginal laceration		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Wound		

subjects affected / exposed occurrences (all)	1 / 113 (0.88%) 1	2 / 108 (1.85%) 2	
Wrist fracture subjects affected / exposed occurrences (all)	1 / 113 (0.88%) 1	0 / 108 (0.00%) 0	
Cardiac disorders			
Bundle branch block right subjects affected / exposed occurrences (all)	0 / 113 (0.00%) 0	1 / 108 (0.93%) 1	
Extrasystoles subjects affected / exposed occurrences (all)	0 / 113 (0.00%) 0	1 / 108 (0.93%) 1	
Palpitations subjects affected / exposed occurrences (all)	3 / 113 (2.65%) 3	2 / 108 (1.85%) 2	
Pericardial effusion subjects affected / exposed occurrences (all)	3 / 113 (2.65%) 3	0 / 108 (0.00%) 0	
Sinus tachycardia subjects affected / exposed occurrences (all)	0 / 113 (0.00%) 0	1 / 108 (0.93%) 1	
Tachycardia subjects affected / exposed occurrences (all)	0 / 113 (0.00%) 0	1 / 108 (0.93%) 4	
Ventricular arrhythmia subjects affected / exposed occurrences (all)	0 / 113 (0.00%) 0	1 / 108 (0.93%) 1	
Nervous system disorders			
Ageusia subjects affected / exposed occurrences (all)	1 / 113 (0.88%) 1	0 / 108 (0.00%) 0	
Burning sensation subjects affected / exposed occurrences (all)	1 / 113 (0.88%) 1	0 / 108 (0.00%) 0	
Cerebral ischaemia			

subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Dizziness		
subjects affected / exposed	14 / 113 (12.39%)	15 / 108 (13.89%)
occurrences (all)	15	23
Dizziness postural		
subjects affected / exposed	5 / 113 (4.42%)	1 / 108 (0.93%)
occurrences (all)	5	1
Dysaesthesia		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Dysgeusia		
subjects affected / exposed	9 / 113 (7.96%)	7 / 108 (6.48%)
occurrences (all)	10	9
Extrapyramidal disorder		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Headache		
subjects affected / exposed	15 / 113 (13.27%)	17 / 108 (15.74%)
occurrences (all)	17	25
Hydrocephalus		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Hypoaesthesia		
subjects affected / exposed	3 / 113 (2.65%)	1 / 108 (0.93%)
occurrences (all)	4	1
Lethargy		
subjects affected / exposed	0 / 113 (0.00%)	2 / 108 (1.85%)
occurrences (all)	0	2
Memory impairment		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	2
Meningism		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Migraine		

subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Motor dysfunction		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Myoclonus		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Neuropathy peripheral		
subjects affected / exposed	8 / 113 (7.08%)	11 / 108 (10.19%)
occurrences (all)	10	15
Paraesthesia		
subjects affected / exposed	9 / 113 (7.96%)	10 / 108 (9.26%)
occurrences (all)	13	13
Parosmia		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Partial seizures		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Peripheral motor neuropathy		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	3	0
Peripheral sensory neuropathy		
subjects affected / exposed	9 / 113 (7.96%)	3 / 108 (2.78%)
occurrences (all)	17	3
Polyneuropathy		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Polyneuropathy in malignant disease		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Restless legs syndrome		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Sciatica		

subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)	
occurrences (all)	1	1	
Syncope			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Tension headache			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Tremor			
subjects affected / exposed	1 / 113 (0.88%)	2 / 108 (1.85%)	
occurrences (all)	1	2	
Blood and lymphatic system disorders			
Anaemia			
subjects affected / exposed	12 / 113 (10.62%)	13 / 108 (12.04%)	
occurrences (all)	25	29	
Anaemia folate deficiency			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Aplastic anaemia			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Haemorrhagic diathesis			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Leukocytosis			
subjects affected / exposed	0 / 113 (0.00%)	2 / 108 (1.85%)	
occurrences (all)	0	3	
Leukopenia			
subjects affected / exposed	8 / 113 (7.08%)	8 / 108 (7.41%)	
occurrences (all)	13	13	
Lymphadenopathy			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Lymphopenia			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	3	0	

Neutropenia			
subjects affected / exposed	16 / 113 (14.16%)	23 / 108 (21.30%)	
occurrences (all)	43	68	
Neutrophilia			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Splenomegaly			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Thrombocytopenia			
subjects affected / exposed	3 / 113 (2.65%)	2 / 108 (1.85%)	
occurrences (all)	17	14	
Thrombocytosis			
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)	
occurrences (all)	2	0	
White blood cell disorder			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Ear and labyrinth disorders			
Deafness			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Ear pain			
subjects affected / exposed	2 / 113 (1.77%)	1 / 108 (0.93%)	
occurrences (all)	2	1	
Hypoacusis			
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)	
occurrences (all)	3	0	
Tinnitus			
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)	
occurrences (all)	2	0	
Vertigo			
subjects affected / exposed	6 / 113 (5.31%)	3 / 108 (2.78%)	
occurrences (all)	6	4	
Eye disorders			

Asthenopia		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Blepharitis		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Dry eye		
subjects affected / exposed	2 / 113 (1.77%)	4 / 108 (3.70%)
occurrences (all)	2	4
Eye discharge		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Eye disorder		
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)
occurrences (all)	2	0
Eye irritation		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	2
Eyelid function disorder		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Eyelid oedema		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Eyelid pain		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Lacrimation increased		
subjects affected / exposed	5 / 113 (4.42%)	3 / 108 (2.78%)
occurrences (all)	7	3
Ocular hyperaemia		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Ocular surface disease		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1

Photopsia			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Scleral discolouration			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Vision blurred			
subjects affected / exposed	7 / 113 (6.19%)	3 / 108 (2.78%)	
occurrences (all)	7	3	
Visual impairment			
subjects affected / exposed	2 / 113 (1.77%)	2 / 108 (1.85%)	
occurrences (all)	2	4	
Vitreous floaters			
subjects affected / exposed	2 / 113 (1.77%)	1 / 108 (0.93%)	
occurrences (all)	2	1	
Gastrointestinal disorders			
Abdominal discomfort			
subjects affected / exposed	3 / 113 (2.65%)	3 / 108 (2.78%)	
occurrences (all)	3	3	
Abdominal distension			
subjects affected / exposed	9 / 113 (7.96%)	10 / 108 (9.26%)	
occurrences (all)	10	11	
Abdominal hernia			
subjects affected / exposed	3 / 113 (2.65%)	0 / 108 (0.00%)	
occurrences (all)	4	0	
Abdominal pain			
subjects affected / exposed	35 / 113 (30.97%)	36 / 108 (33.33%)	
occurrences (all)	58	67	
Abdominal pain lower			
subjects affected / exposed	5 / 113 (4.42%)	7 / 108 (6.48%)	
occurrences (all)	6	9	
Abdominal pain upper			
subjects affected / exposed	18 / 113 (15.93%)	16 / 108 (14.81%)	
occurrences (all)	21	25	
Abdominal tenderness			



subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	1
Abnormal faeces		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Anal hypoaesthesia		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	2	0
Anal incontinence		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	1
Anorectal discomfort		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Aphthous ulcer		
subjects affected / exposed	6 / 113 (5.31%)	2 / 108 (1.85%)
occurrences (all)	9	4
Ascites		
subjects affected / exposed	27 / 113 (23.89%)	8 / 108 (7.41%)
occurrences (all)	57	16
Cheilitis		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Constipation		
subjects affected / exposed	39 / 113 (34.51%)	36 / 108 (33.33%)
occurrences (all)	63	66
Diarrhoea		
subjects affected / exposed	32 / 113 (28.32%)	28 / 108 (25.93%)
occurrences (all)	62	43
Diverticulum intestinal		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Dry mouth		
subjects affected / exposed	6 / 113 (5.31%)	5 / 108 (4.63%)
occurrences (all)	7	5
Dyschezia		

subjects affected / exposed	3 / 113 (2.65%)	0 / 108 (0.00%)
occurrences (all)	3	0
Dyspepsia		
subjects affected / exposed	21 / 113 (18.58%)	15 / 108 (13.89%)
occurrences (all)	26	17
Dysphagia		
subjects affected / exposed	6 / 113 (5.31%)	5 / 108 (4.63%)
occurrences (all)	6	5
Enteritis		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Epigastric discomfort		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Eructation		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	2	1
Flatulence		
subjects affected / exposed	3 / 113 (2.65%)	3 / 108 (2.78%)
occurrences (all)	3	3
Gastric disorder		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Gastric stenosis		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Gastritis		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Gastritis erosive		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Gastrointestinal motility disorder		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Gastrointestinal pain		

subjects affected / exposed	2 / 113 (1.77%)	1 / 108 (0.93%)
occurrences (all)	4	2
Gastrointestinal sounds abnormal		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Gastrooesophageal reflux disease		
subjects affected / exposed	4 / 113 (3.54%)	8 / 108 (7.41%)
occurrences (all)	4	9
Gingival bleeding		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Gingival pain		
subjects affected / exposed	4 / 113 (3.54%)	3 / 108 (2.78%)
occurrences (all)	4	3
Glossodynia		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Haematemesis		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Haematochezia		
subjects affected / exposed	1 / 113 (0.88%)	2 / 108 (1.85%)
occurrences (all)	1	2
Haemorrhoidal haemorrhage		
subjects affected / exposed	0 / 113 (0.00%)	2 / 108 (1.85%)
occurrences (all)	0	2
Haemorrhoids		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	1
Hiatus hernia		
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)
occurrences (all)	2	0
Hyperchlorhydria		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Ileus		

subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Intestinal obstruction		
subjects affected / exposed	0 / 113 (0.00%)	2 / 108 (1.85%)
occurrences (all)	0	2
Intra-abdominal fluid collection		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	1
Mouth ulceration		
subjects affected / exposed	4 / 113 (3.54%)	6 / 108 (5.56%)
occurrences (all)	6	13
Nausea		
subjects affected / exposed	66 / 113 (58.41%)	61 / 108 (56.48%)
occurrences (all)	134	113
Odynophagia		
subjects affected / exposed	1 / 113 (0.88%)	3 / 108 (2.78%)
occurrences (all)	2	3
Oesophagitis		
subjects affected / exposed	3 / 113 (2.65%)	1 / 108 (0.93%)
occurrences (all)	3	1
Oral dysaesthesia		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Oral pain		
subjects affected / exposed	2 / 113 (1.77%)	6 / 108 (5.56%)
occurrences (all)	2	7
Paraesthesia oral		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Periodontal disease		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Proctalgia		
subjects affected / exposed	2 / 113 (1.77%)	2 / 108 (1.85%)
occurrences (all)	4	3
Rectal discharge		

subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)	
occurrences (all)	1	1	
Rectal haemorrhage			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Reflux gastritis			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Small intestinal obstruction			
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)	
occurrences (all)	2	0	
Stomatitis			
subjects affected / exposed	58 / 113 (51.33%)	55 / 108 (50.93%)	
occurrences (all)	112	137	
Swollen tongue			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Tongue coated			
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)	
occurrences (all)	2	0	
Tongue disorder			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Tooth discolouration			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Tooth loss			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Toothache			
subjects affected / exposed	1 / 113 (0.88%)	3 / 108 (2.78%)	
occurrences (all)	1	4	
Vomiting			
subjects affected / exposed	49 / 113 (43.36%)	35 / 108 (32.41%)	
occurrences (all)	76	75	
Skin and subcutaneous tissue disorders			

Acanthosis nigricans		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	2	1
Acne		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Blister		
subjects affected / exposed	7 / 113 (6.19%)	4 / 108 (3.70%)
occurrences (all)	10	5
Alopecia		
subjects affected / exposed	21 / 113 (18.58%)	12 / 108 (11.11%)
occurrences (all)	27	15
Dermatitis		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Decubitus ulcer		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	2	0
Dermatitis acneiform		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Dermatitis allergic		
subjects affected / exposed	3 / 113 (2.65%)	2 / 108 (1.85%)
occurrences (all)	8	2
Dermatitis exfoliative		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	2
Eczema		
subjects affected / exposed	3 / 113 (2.65%)	1 / 108 (0.93%)
occurrences (all)	3	2
Dry skin		
subjects affected / exposed	10 / 113 (8.85%)	13 / 108 (12.04%)
occurrences (all)	11	13
Erythema		
subjects affected / exposed	3 / 113 (2.65%)	5 / 108 (4.63%)
occurrences (all)	9	10

Exfoliative rash		
subjects affected / exposed	4 / 113 (3.54%)	1 / 108 (0.93%)
occurrences (all)	6	1
Generalised erythema		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Hyperhidrosis		
subjects affected / exposed	1 / 113 (0.88%)	2 / 108 (1.85%)
occurrences (all)	1	3
Nail discolouration		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	1
Nail disorder		
subjects affected / exposed	2 / 113 (1.77%)	4 / 108 (3.70%)
occurrences (all)	2	4
Nail dystrophy		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Night sweats		
subjects affected / exposed	4 / 113 (3.54%)	2 / 108 (1.85%)
occurrences (all)	4	4
Pain of skin		
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)
occurrences (all)	2	0
Onycholysis		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Palmar erythema		
subjects affected / exposed	1 / 113 (0.88%)	2 / 108 (1.85%)
occurrences (all)	1	2
Panniculitis		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Palmar-plantar erythrodysesthesia syndrome		

subjects affected / exposed	70 / 113 (61.95%)	61 / 108 (56.48%)
occurrences (all)	247	177
Plantar erythema		
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)
occurrences (all)	2	0
Pigmentation disorder		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	2	0
Pruritus		
subjects affected / exposed	11 / 113 (9.73%)	8 / 108 (7.41%)
occurrences (all)	17	10
Pruritus generalised		
subjects affected / exposed	3 / 113 (2.65%)	1 / 108 (0.93%)
occurrences (all)	5	1
Psoriasis		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	2	0
Rash		
subjects affected / exposed	30 / 113 (26.55%)	28 / 108 (25.93%)
occurrences (all)	59	67
Pustular psoriasis		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Rash erythematous		
subjects affected / exposed	2 / 113 (1.77%)	3 / 108 (2.78%)
occurrences (all)	2	3
Rash generalised		
subjects affected / exposed	3 / 113 (2.65%)	0 / 108 (0.00%)
occurrences (all)	4	0
Rash macular		
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)
occurrences (all)	2	0
Rash maculo-papular		
subjects affected / exposed	4 / 113 (3.54%)	4 / 108 (3.70%)
occurrences (all)	6	26
Rash pruritic		



subjects affected / exposed	2 / 113 (1.77%)	2 / 108 (1.85%)
occurrences (all)	2	2
Rash vesicular		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Scar pain		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	1
Skin discolouration		
subjects affected / exposed	2 / 113 (1.77%)	2 / 108 (1.85%)
occurrences (all)	2	2
Skin disorder		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Skin exfoliation		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	4	1
Skin fissures		
subjects affected / exposed	2 / 113 (1.77%)	1 / 108 (0.93%)
occurrences (all)	2	1
Skin hyperpigmentation		
subjects affected / exposed	7 / 113 (6.19%)	11 / 108 (10.19%)
occurrences (all)	8	14
Skin irritation		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Skin lesion		
subjects affected / exposed	0 / 113 (0.00%)	3 / 108 (2.78%)
occurrences (all)	0	3
Skin reaction		
subjects affected / exposed	0 / 113 (0.00%)	2 / 108 (1.85%)
occurrences (all)	0	4
Skin mass		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Skin tightness		

subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Skin ulcer			
subjects affected / exposed	0 / 113 (0.00%)	2 / 108 (1.85%)	
occurrences (all)	0	2	
Stasis dermatitis			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Renal and urinary disorders			
Anuria			
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)	
occurrences (all)	2	0	
Dysuria			
subjects affected / exposed	6 / 113 (5.31%)	5 / 108 (4.63%)	
occurrences (all)	6	6	
Haematuria			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	2	
Hydronephrosis			
subjects affected / exposed	1 / 113 (0.88%)	2 / 108 (1.85%)	
occurrences (all)	1	2	
Kidney enlargement			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Nocturia			
subjects affected / exposed	1 / 113 (0.88%)	2 / 108 (1.85%)	
occurrences (all)	1	2	
Obstructive uropathy			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Oliguria			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Pollakiuria			
subjects affected / exposed	3 / 113 (2.65%)	1 / 108 (0.93%)	
occurrences (all)	3	1	

Proteinuria			
subjects affected / exposed	6 / 113 (5.31%)	4 / 108 (3.70%)	
occurrences (all)	6	10	
Renal colic			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Renal pain			
subjects affected / exposed	0 / 113 (0.00%)	2 / 108 (1.85%)	
occurrences (all)	0	2	
Renal vein thrombosis			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Urinary hesitation			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Urinary incontinence			
subjects affected / exposed	3 / 113 (2.65%)	2 / 108 (1.85%)	
occurrences (all)	3	2	
Urinary retention			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Urinary tract obstruction			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Endocrine disorders			
Cushingoid			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Hypothyroidism			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Musculoskeletal and connective tissue disorders			
Arthralgia			
subjects affected / exposed	11 / 113 (9.73%)	7 / 108 (6.48%)	
occurrences (all)	18	8	
Arthritis			

subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Back pain		
subjects affected / exposed	12 / 113 (10.62%)	17 / 108 (15.74%)
occurrences (all)	13	20
Bone pain		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	1
Bursitis		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Coccydynia		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Enthesopathy		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Flank pain		
subjects affected / exposed	3 / 113 (2.65%)	4 / 108 (3.70%)
occurrences (all)	5	6
Fracture pain		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Groin pain		
subjects affected / exposed	2 / 113 (1.77%)	2 / 108 (1.85%)
occurrences (all)	3	2
Intervertebral disc displacement		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Joint swelling		
subjects affected / exposed	0 / 113 (0.00%)	2 / 108 (1.85%)
occurrences (all)	0	2
Limb discomfort		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Muscle fatigue		

subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Muscle spasms		
subjects affected / exposed	9 / 113 (7.96%)	11 / 108 (10.19%)
occurrences (all)	10	20
Muscle tightness		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Muscular weakness		
subjects affected / exposed	0 / 113 (0.00%)	3 / 108 (2.78%)
occurrences (all)	0	3
Musculoskeletal chest pain		
subjects affected / exposed	3 / 113 (2.65%)	8 / 108 (7.41%)
occurrences (all)	4	9
Musculoskeletal discomfort		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Musculoskeletal disorder		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Musculoskeletal pain		
subjects affected / exposed	4 / 113 (3.54%)	7 / 108 (6.48%)
occurrences (all)	4	9
Musculoskeletal stiffness		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	1
Myalgia		
subjects affected / exposed	4 / 113 (3.54%)	2 / 108 (1.85%)
occurrences (all)	4	3
Neck pain		
subjects affected / exposed	4 / 113 (3.54%)	3 / 108 (2.78%)
occurrences (all)	4	3
Osteopenia		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Pain in extremity		

subjects affected / exposed	13 / 113 (11.50%)	4 / 108 (3.70%)	
occurrences (all)	17	4	
Spinal osteoarthritis			
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)	
occurrences (all)	1	1	
Spinal pain			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Temporomandibular joint syndrome			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Tendon disorder			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Trismus			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Infections and infestations			
Angular cheilitis			
subjects affected / exposed	1 / 113 (0.88%)	2 / 108 (1.85%)	
occurrences (all)	1	2	
Appendicitis			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Bronchitis			
subjects affected / exposed	2 / 113 (1.77%)	3 / 108 (2.78%)	
occurrences (all)	2	3	
Candida infection			
subjects affected / exposed	2 / 113 (1.77%)	3 / 108 (2.78%)	
occurrences (all)	3	3	
Carbuncle			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Cellulitis			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	

Conjunctivitis		
subjects affected / exposed	4 / 113 (3.54%)	3 / 108 (2.78%)
occurrences (all)	4	3
Cystitis		
subjects affected / exposed	3 / 113 (2.65%)	3 / 108 (2.78%)
occurrences (all)	6	3
Device related infection		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Ear infection		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Enterocolitis infectious		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Erysipelas		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Escherichia urinary tract infection		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Eye infection		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	1
Folliculitis		
subjects affected / exposed	3 / 113 (2.65%)	4 / 108 (3.70%)
occurrences (all)	5	5
Fungal cystitis		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Fungal infection		
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)
occurrences (all)	2	0
Fungal skin infection		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1

Gastroenteritis		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	1
Gastroenteritis viral		
subjects affected / exposed	0 / 113 (0.00%)	2 / 108 (1.85%)
occurrences (all)	0	5
Gastrointestinal infection		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Gingivitis		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Herpes virus infection		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	1
Herpes zoster		
subjects affected / exposed	2 / 113 (1.77%)	1 / 108 (0.93%)
occurrences (all)	2	1
Hordeolum		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Infection		
subjects affected / exposed	1 / 113 (0.88%)	2 / 108 (1.85%)
occurrences (all)	2	2
Influenza		
subjects affected / exposed	1 / 113 (0.88%)	4 / 108 (3.70%)
occurrences (all)	1	4
Lice infestation		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Localised infection		
subjects affected / exposed	2 / 113 (1.77%)	1 / 108 (0.93%)
occurrences (all)	3	2
Lower respiratory tract infection		
subjects affected / exposed	2 / 113 (1.77%)	4 / 108 (3.70%)
occurrences (all)	3	4



Lung abscess		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Lung infection		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Lymphangitis		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Mastoiditis		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Nail infection		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	1
Nasopharyngitis		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Oesophageal candidiasis		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Onychomycosis		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Oral candidiasis		
subjects affected / exposed	6 / 113 (5.31%)	9 / 108 (8.33%)
occurrences (all)	6	10
Oral fungal infection		
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)
occurrences (all)	2	0
Oral herpes		
subjects affected / exposed	3 / 113 (2.65%)	1 / 108 (0.93%)
occurrences (all)	3	1
Oropharyngeal candidiasis		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0

Paronychia		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	3
Periorbital cellulitis		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Pharyngitis		
subjects affected / exposed	3 / 113 (2.65%)	3 / 108 (2.78%)
occurrences (all)	3	3
Pneumonia		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	1
Post procedural cellulitis		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Pseudomonas infection		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	1
Pyuria		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	1
Rash pustular		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Respiratory tract infection		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Rhinitis		
subjects affected / exposed	4 / 113 (3.54%)	4 / 108 (3.70%)
occurrences (all)	6	5
Skin infection		
subjects affected / exposed	0 / 113 (0.00%)	2 / 108 (1.85%)
occurrences (all)	0	2
Soft tissue infection		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0

Staphylococcal infection		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	2
Tinea infection		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Tonsillitis		
subjects affected / exposed	0 / 113 (0.00%)	2 / 108 (1.85%)
occurrences (all)	0	2
Tooth abscess		
subjects affected / exposed	1 / 113 (0.88%)	2 / 108 (1.85%)
occurrences (all)	1	2
Tooth infection		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	1	1
Upper respiratory tract infection		
subjects affected / exposed	12 / 113 (10.62%)	7 / 108 (6.48%)
occurrences (all)	13	10
Upper respiratory tract infection bacterial		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Urinary tract infection		
subjects affected / exposed	7 / 113 (6.19%)	9 / 108 (8.33%)
occurrences (all)	7	14
Urinary tract infection bacterial		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Viral infection		
subjects affected / exposed	1 / 113 (0.88%)	1 / 108 (0.93%)
occurrences (all)	2	1
Viral pharyngitis		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Viral upper respiratory tract infection		

subjects affected / exposed	11 / 113 (9.73%)	11 / 108 (10.19%)	
occurrences (all)	14	13	
Vulvovaginal candidiasis			
subjects affected / exposed	3 / 113 (2.65%)	3 / 108 (2.78%)	
occurrences (all)	3	3	
Wound infection			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Metabolism and nutrition disorders			
Cachexia			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	3	0	
Decreased appetite			
subjects affected / exposed	27 / 113 (23.89%)	24 / 108 (22.22%)	
occurrences (all)	33	36	
Dehydration			
subjects affected / exposed	2 / 113 (1.77%)	4 / 108 (3.70%)	
occurrences (all)	2	4	
Diabetes mellitus			
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)	
occurrences (all)	1	0	
Glucose tolerance impaired			
subjects affected / exposed	0 / 113 (0.00%)	2 / 108 (1.85%)	
occurrences (all)	0	2	
Hypercalcaemia			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	
Hyperglycaemia			
subjects affected / exposed	3 / 113 (2.65%)	1 / 108 (0.93%)	
occurrences (all)	8	1	
Hyperkalaemia			
subjects affected / exposed	2 / 113 (1.77%)	0 / 108 (0.00%)	
occurrences (all)	4	0	
Hypermagnesaemia			
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)	
occurrences (all)	0	1	

Hyperuricaemia		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Hypoalbuminaemia		
subjects affected / exposed	4 / 113 (3.54%)	3 / 108 (2.78%)
occurrences (all)	9	3
Hypocalcaemia		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Hypochloraemia		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Hypoglycaemia		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	3	0
Hypokalaemia		
subjects affected / exposed	23 / 113 (20.35%)	11 / 108 (10.19%)
occurrences (all)	37	14
Hypomagnesaemia		
subjects affected / exposed	11 / 113 (9.73%)	3 / 108 (2.78%)
occurrences (all)	25	7
Hyponatraemia		
subjects affected / exposed	4 / 113 (3.54%)	6 / 108 (5.56%)
occurrences (all)	4	9
Hypophosphataemia		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1
Hypoproteinaemia		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	1	0
Hypovolaemia		
subjects affected / exposed	1 / 113 (0.88%)	0 / 108 (0.00%)
occurrences (all)	2	0
Iron deficiency		
subjects affected / exposed	0 / 113 (0.00%)	1 / 108 (0.93%)
occurrences (all)	0	1



## More information

### Substantial protocol amendments (globally)

Were there any global substantial amendments to the protocol? Yes

Date	Amendment
31 October 2012	<ul style="list-style-type: none"><li>- To provide current language for the management of trebananib related toxicities. Pleural effusion and ascites are known identified risks of trebananib, and thus a new toxicity management section was added to provide guidance to participating investigators of the study.</li><li>- Statistical changes rationale are as follows:<ul style="list-style-type: none"><li>- The accrual assumptions for this study were modified to reflect the initial accrual of 58 subjects and the accrual hold. Accrual is most commonly modeled with a 2-stage accrual curve, with 50% of the accrual occurring in the last 30% of the accrual time. To reflect the initial accrual, the accrual hold, and the future accrual for this study, accrual is modeled with a 4-part accrual curve:<ul style="list-style-type: none"><li>-- Part 1 reflects linear accrual of the first 58 patients over 7 months</li><li>-- Part 2 reflects the hold of 15 months</li><li>-- Parts 3 and 4 are the typical 2-stage accrual used; part 3 reflects accrual of the remaining 322 patients, with 47% of this accrual occurring over the 16 months and the remaining 53% of accrual occurring over the last 6 months.</li></ul></li><li>- the operating characteristics of this futility analysis are based on the hypothesis testing being set up as 1-sided 2.5% testing. Therefore, the references to 2-sided 5% tests have been changed to 1-sided 2.5% tests throughout.</li><li>- The primary analysis was modified to reflect the inclusion of the time of accrual (before or after the accrual hold) as a stratification factor</li><li>- To reword and move hypertension from exclusion to inclusion criteria</li><li>- To add guidelines regarding immune modulators as excluded medications during the study</li><li>- To add the new pregnancy and lactation reporting guidance</li><li>- To update language on the reporting of adverse events by Investigators and the use of Appendix B in the Investigator's Brochure for the determination of adverse event expectedness</li><li>- Update the trebananib Investigator's Brochure version number and date</li><li>- Administrative and typographical corrections throughout</li></ul></li></ul>
23 January 2014	<ul style="list-style-type: none"><li>- To address the decisions to close the study to further randomization last 23 October 2013 due to global supply shortages of DOXIL® (doxorubicin HCl liposome injection)/CAELYX® (pegylated liposomal doxorubicin hydrochloride)<ul style="list-style-type: none"><li>- A total of 223 subjects were randomized to the study</li><li>- Statistical analyses plans were amended to reflect the change in the number of subjects on study</li></ul></li><li>- To add language on how to address and report serious adverse events occurring outside the protocol required reporting period and provide serious adverse events sample forms</li><li>- Update key contact information of the Clinical Research Medical Director (Medical Monitor) and study management for the study</li><li>- Update toxicity management guidelines</li><li>- Update the references in Section 13</li><li>- Correct any typographical and/or grammatical errors in the protocol</li></ul>

Notes:

### Interruptions (globally)

Were there any global interruptions to the trial? Yes

Date	Interruption	Restart date
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23 November 2011	Due to a global shortage of PLD, enrollment was put on hold from 23 November 2011 (last subject enrolled date pre-hold) until 10 January 2013 (first subjected enrolled date post-hold).	10 January 2013
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Notes:

## Limitations and caveats

None reported