

**Clinical trial results:****An Open-Label, Multicenter Study to Assess the Safety of RO5185426 (Vemurafenib) in Patients with Metastatic Melanoma****Summary**

EudraCT number	2010-023526-21
Trial protocol	ES AT SE NL SI GR DE GB CZ HU SK BE BG NO PT LV FI IT DK
Global end of trial date	54 February 2016

**Results information**

Result version number	v1 (current)
This version publication date	06 August 2017
First version publication date	06 August 2017

**Trial information****Trial identification**

Sponsor protocol code	MO25515
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**Additional study identifiers**

ISRCTN number	-
ClinicalTrials.gov id (NCT number)	NCT01307397
WHO universal trial number (UTN)	-

Notes:

**Sponsors**

Sponsor organisation name	Hoffmann-La Roche
Sponsor organisation address	Grenzacherstrasse, Basel, Switzerland, CH-4070
Public contact	Roche Trial Information Hotline, F. Hoffmann-La Roche AG, +41 61 6878333, global.trial_information@roche.com
Scientific contact	Roche Trial Information Hotline, F. Hoffmann-La Roche AG, +41 61 6878333, global.trial_information@roche.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?	Yes

Notes:

## Results analysis stage

Analysis stage	Final
Date of interim/final analysis	14 March 2016
Is this the analysis of the primary completion data?	No
Global end of trial reached?	Yes
Global end of trial date	24 February 2016
Was the trial ended prematurely?	No

Notes:

## General information about the trial

Main objective of the trial:

To evaluate the safety and tolerability of vemurafenib in participants with metastatic melanoma (surgically incurable and unresectable Stage IIIC or Stage IV, American Joint Committee on Cancer [AJCC]), harboring the BRAF V600 mutation (identified by the cobas® 4800 BRAF V600 Mutation Test)

Protection of trial subjects:

The study was conducted in accordance with the principles of the "Declaration of Helsinki" and Good Clinical Practice (GCP) according to the regulations and procedures described in the protocol. Approval from the Ethics Committee (EC)/Institutional Review Board (IRB) was obtained before study start. Roche also obtained approval from the relevant Competent Authority prior to starting the study. No modifications were made to the protocol after receipt of the EC/IRB approval.

Background therapy: -

Evidence for comparator: -

Actual start date of recruitment	01 March 2011
Long term follow-up planned	No
Independent data monitoring committee (IDMC) involvement?	No

Notes:

## Population of trial subjects

### Subjects enrolled per country

Country: Number of subjects enrolled	Albania: 12
Country: Number of subjects enrolled	Argentina: 19
Country: Number of subjects enrolled	Australia: 158
Country: Number of subjects enrolled	Austria: 59
Country: Number of subjects enrolled	Belgium: 66
Country: Number of subjects enrolled	Bosnia and Herzegovina: 27
Country: Number of subjects enrolled	Brazil: 53
Country: Number of subjects enrolled	Bulgaria: 42
Country: Number of subjects enrolled	Canada: 145
Country: Number of subjects enrolled	Colombia: 17
Country: Number of subjects enrolled	Croatia: 21
Country: Number of subjects enrolled	Czech Republic: 137
Country: Number of subjects enrolled	Denmark: 28
Country: Number of subjects enrolled	Ecuador: 1
Country: Number of subjects enrolled	Estonia: 8
Country: Number of subjects enrolled	Finland: 18
Country: Number of subjects enrolled	Germany: 389
Country: Number of subjects enrolled	Greece: 55

Country: Number of subjects enrolled	Hungary: 55
Country: Number of subjects enrolled	India: 3
Country: Number of subjects enrolled	Ireland: 12
Country: Number of subjects enrolled	Israel: 73
Country: Number of subjects enrolled	Italy: 383
Country: Number of subjects enrolled	Korea, Republic of: 41
Country: Number of subjects enrolled	Latvia: 12
Country: Number of subjects enrolled	Lithuania: 13
Country: Number of subjects enrolled	Macedonia, the former Yugoslav Republic of: 20
Country: Number of subjects enrolled	Mexico: 10
Country: Number of subjects enrolled	Netherlands: 234
Country: Number of subjects enrolled	Norway: 47
Country: Number of subjects enrolled	Peru: 5
Country: Number of subjects enrolled	Poland: 75
Country: Number of subjects enrolled	Portugal: 21
Country: Number of subjects enrolled	Romania: 29
Country: Number of subjects enrolled	Russian Federation: 60
Country: Number of subjects enrolled	Serbia: 28
Country: Number of subjects enrolled	Slovakia: 18
Country: Number of subjects enrolled	Slovenia: 21
Country: Number of subjects enrolled	South Africa: 34
Country: Number of subjects enrolled	Spain: 300
Country: Number of subjects enrolled	Sweden: 77
Country: Number of subjects enrolled	Switzerland: 36
Country: Number of subjects enrolled	Turkey: 111
Country: Number of subjects enrolled	United Kingdom: 246
Worldwide total number of subjects	3219
EEA total number of subjects	2366

Notes:

### Subjects enrolled per age group

In utero	0
Preterm newborn - gestational age < 37 wk	0
Newborns (0-27 days)	0
Infants and toddlers (28 days-23 months)	0
Children (2-11 years)	0
Adolescents (12-17 years)	3
Adults (18-64 years)	2371
From 65 to 84 years	825
85 years and over	20

## Subject disposition

### Recruitment

Recruitment details: -

### Pre-assignment

Screening details:

A total of 7495 participants were screened; 3224 participants were enrolled and 3219 participants received at least 1 dose of vemurafenib. Further 38 participants were screened at a site in Austria and 29 of these participants were enrolled and treated; however, they were not included in analysis populations due to critical audit findings.

### Period 1

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

### Arms

<b>Arm title</b>	Vemurafenib
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Arm description:

Participants received continuous oral doses of vemurafenib 960 milligrams (mg) (four 240 mg tablets) twice daily in each 28-day treatment cycle until the development of progressive disease, unacceptable toxicity, consent withdrawal, protocol violations endangering participant's safety, death or study termination by the Sponsor.

Arm type	Experimental
Investigational medicinal product name	Vemurafenib
Investigational medicinal product code	
Other name	RO5185426, Zelboraf
Pharmaceutical forms	Film-coated tablet
Routes of administration	Oral use

Dosage and administration details:

Participants received continuous oral doses of vemurafenib 960 mg (four 240 mg tablets) twice daily in each 28-day treatment cycle until the development of progressive disease, unacceptable toxicity, consent withdrawal, protocol violations endangering participant's safety, death or study termination by the Sponsor.

<b>Number of subjects in period 1</b>	Vemurafenib
Started	3219
Completed	0
Not completed	3219
Consent withdrawn by subject	62
Investigator's Decision	62
Death	120
Refused Treatment	66
Progressive Disease	2414
Administrative/Other	226
Adverse Events	225

Lost to follow-up	34
Missing	4
Protocol deviation	6

## Baseline characteristics

### Reporting groups

Reporting group title	Vemurafenib
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Reporting group description:

Participants received continuous oral doses of vemurafenib 960 milligrams (mg) (four 240 mg tablets) twice daily in each 28-day treatment cycle until the development of progressive disease, unacceptable toxicity, consent withdrawal, protocol violations endangering participant's safety, death or study termination by the Sponsor.

Reporting group values	Vemurafenib	Total	
Number of subjects	3219	3219	
Age categorical Units: Subjects			
Age Continuous			
The safety population included all participants who received at least one dose of study medication			
Units: years arithmetic mean standard deviation	54.5 ± 14.06	-	
Gender, Male/Female Units: Subjects			
Female	1397	1397	
Male	1822	1822	

## End points

### End points reporting groups

Reporting group title	Vemurafenib
Reporting group description: Participants received continuous oral doses of vemurafenib 960 milligrams (mg) (four 240 mg tablets) twice daily in each 28-day treatment cycle until the development of progressive disease, unacceptable toxicity, consent withdrawal, protocol violations endangering participant's safety, death or study termination by the Sponsor.	

### Primary: Percentage of Participants Experiencing Any Grade 3 or 4 Adverse Events (AEs) as determined by National Cancer Institute-Common Toxicity Criteria for Adverse Events (NCI-CTCAE) Version 4.0

End point title	Percentage of Participants Experiencing Any Grade 3 or 4 Adverse Events (AEs) as determined by National Cancer Institute-Common Toxicity Criteria for Adverse Events (NCI-CTCAE) Version 4.0 <sup>[1]</sup>
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#### End point description:

The intensity of AEs were graded on a 5-point scale (Grade 1 to 5) according to the NCI-CTCAE version 4.0, where Grade 1 indicates "Mild" severity and Grade 5 indicates "Death". The CTCAE defines Grades 3 and 4 as follows: Grade 3 means "Severe"; Inability to work or perform normal daily activity; treatment or medical intervention is indicated in order to improve the overall wellbeing or symptoms; delaying the onset of treatment is not putting the survival of the participant at direct risk. Grade 4 means "Life-threatening, Disabling"; based on extreme limitation in activity; significant medical intervention/therapy required; and hospitalization probable. Analysis was performed on the safety population (all participants who received at least one dose of study medication).

End point type	Primary
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#### End point timeframe:

Baseline up to 28 days post end of treatment (maximum up to 46 months)

#### Notes:

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

Justification: No hypothesis testing was done in this single arm Safety study. All results presented are descriptive only.

<b>End point values</b>	Vemurafenib			
Subject group type	Reporting group			
Number of subjects analysed	3219			
Units: Percentage of participants				
number (not applicable)	52.8			

### Statistical analyses

No statistical analyses for this end point

### Primary: Percentage of Participants With at Least 1 AE Leading to Study Drug Interruption or Drug Discontinuation

End point title	Percentage of Participants With at Least 1 AE Leading to Study Drug Interruption or Drug Discontinuation <sup>[2]</sup>
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End point description:

An AE was considered as any unfavorable and unintended sign, symptom, or disease associated with the use of the study drug, whether or not considered related to the study drug. Pre existing conditions that worsened during the study and laboratory or clinical tests that resulted in a change in treatment or discontinuation from study drug were reported as adverse events. Percentage of participants with dose interruption or discontinuation due to AE was presented. Analysis was performed on the safety population.

End point type | Primary

End point timeframe:

Baseline up to 28 days post end of treatment (maximum up to 46 months)

Notes:

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

Justification: No hypothesis testing was done in this single arm Safety study. All results presented are descriptive only.

End point values	Vemurafenib			
Subject group type	Reporting group			
Number of subjects analysed	3219			
Units: Percentage of participants				
number (not applicable)				
AE Leading to Drug Discontinuation	7			
AE Leading to Study Drug Interruption	34			

## Statistical analyses

No statistical analyses for this end point

## Primary: Percentage of Participants with AEs of Special Interest

End point title | Percentage of Participants with AEs of Special Interest<sup>[3]</sup>

End point description:

AEs of special interest included cutaneous squamous cell carcinoma (SCC), rash, photosensitivity, liver injury, arthralgia, fatigue, gastrointestinal (GI) polyps, pancreatitis, Potentiation of radiation toxicity, prolongation of cardiac repolarization or arrhythmia, non-cutaneous SCC and other primary malignancies (other than cutaneous SCC or new primary melanoma). Analysis was performed on the safety population.

End point type | Primary

End point timeframe:

Baseline up to 28 days post end of treatment (maximum up to 46 months)

Notes:

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

Justification: No hypothesis testing was done in this single arm Safety study. All results presented are descriptive only.

End point values	Vemurafenib			
Subject group type	Reporting group			
Number of subjects analysed	3219			
Units: Percentage of participants				
number (not applicable)				
Arthralgia	42.3			
Rash	47.9			
Photosensitivity	28.4			

Fatigue	36.8			
Cutaneous SCC	14.6			
Non Cutaneous SCC	0.1			
New Primary Melanoma	1.7			
GI Polyps	0.03			
Pancreatitis	0.2			
Potential of radiation toxicity	1.4			
Prolongation of Cardiac Repolarization/ Arrhythmia	17			
Other primary malignancy	3.2			
Liver Injury	14.1			

## Statistical analyses

No statistical analyses for this end point

### Primary: Mean Cumulative Dose of Vemurafenib

End point title	Mean Cumulative Dose of Vemurafenib <sup>[4]</sup>
End point description:	Analysis was performed on the safety population.
End point type	Primary
End point timeframe:	Baseline up to end of treatment or death (maximum up to 46 months)

Notes:

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

Justification: No hypothesis testing was done in this single arm Safety study. All results presented are descriptive only.

<b>End point values</b>	Vemurafenib			
Subject group type	Reporting group			
Number of subjects analysed	3219			
Units: Grams				
arithmetic mean (standard deviation)	501.283 (± 510.9121)			

## Statistical analyses

No statistical analyses for this end point

### Primary: Duration of Vemurafenib Treatment

End point title	Duration of Vemurafenib Treatment <sup>[5]</sup>
End point description:	Exposure excluding treatment interruptions: Duration during which participants actually took vemurafenib. Any time without dose taken due to adverse events, noncompliance or any other reasons was not counted. Exposure including treatment interruptions: date of last dose date of first dose + 1; duration during which participants actually took vemurafenib as well as duration on which medication was not taken were included in this calculation. Analysis was performed on the safety population.
End point type	Primary

End point timeframe:

Baseline up to end of treatment or death (maximum up to 46 months)

Notes:

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

Justification: No hypothesis testing was done in this single arm Safety study. All results presented are descriptive only.

<b>End point values</b>	Vemurafenib			
Subject group type	Reporting group			
Number of subjects analysed	3219			
Units: Months				
arithmetic mean (standard deviation)				
Exposure excluding interruptions	9.383 ( $\pm$ 9.6755)			
Exposure including interruptions	9.724 ( $\pm$ 9.9072)			

## Statistical analyses

No statistical analyses for this end point

## Primary: Mean Total Vemurafenib Dose Per Day

End point title | Mean Total Vemurafenib Dose Per Day<sup>[6]</sup>

End point description:

Exposure excluding treatment interruptions: Duration during which participants actually took vemurafenib. Any time without dose taken due to adverse events, noncompliance or any other reasons was not counted. Exposure including treatment interruptions: date of last dose date of first dose + 1; duration during which participants actually took vemurafenib as well as duration on which medication was not taken were included in this calculation. Average total dose per day: total actual dose taken divided by total actual days on treatment. Analysis was performed on the safety population.

End point type | Primary

End point timeframe:

Baseline up to end of treatment or death (maximum up to 46 months)

Notes:

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

Justification: No hypothesis testing was done in this single arm Safety study. All results presented are descriptive only.

<b>End point values</b>	Vemurafenib			
Subject group type	Reporting group			
Number of subjects analysed	3219			
Units: Grams				
arithmetic mean (standard deviation)				
Average Total Dose Per Day Excluding Interruptions	1.802 ( $\pm$ 0.2314)			
Average Total Dose Per Day Including Interruptions	1.732 ( $\pm$ 0.2874)			

## Statistical analyses

No statistical analyses for this end point

### Primary: Dose Intensity of vemurafenib

End point title	Dose Intensity of vemurafenib <sup>[7]</sup>
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End point description:

Dose intensity was defined as (total actual doses taken/total planned doses) \*100, where total planned doses = prescribed doses \* planned days on treatment, where planned days on treatment were defined as the interval between date of first dose and date of last dose. Analysis was performed on the safety population.

End point type	Primary
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End point timeframe:

Baseline up to end of treatment or death (maximum up to 46 months)

Notes:

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

Justification: No hypothesis testing was done in this single arm Safety study. All results presented are descriptive only.

End point values	Vemurafenib			
Subject group type	Reporting group			
Number of subjects analysed	3219			
Units: Percentage of planned dose				
arithmetic mean (standard deviation)	90.21 (± 14.966)			

## Statistical analyses

No statistical analyses for this end point

### Secondary: Percentage of Participants With Improvement in Eastern Cooperative Group (ECOG) Performance Status

End point title	Percentage of Participants With Improvement in Eastern Cooperative Group (ECOG) Performance Status
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End point description:

ECOG Performance Status was measured on therapy assessed participant's performance status on 5 point scale: 0 = fully active/able to carry on all pre disease activities without restriction; 1=restricted in physically strenuous activity, ambulatory/able to carry out light or sedentary work; 2=ambulatory (greater than [ $>$ ] 50% of waking hours [hrs]), capable of all self care, unable to carry out any work activities; 3=capable of only limited self care, confined to bed/chair  $>$ 50% of waking hrs; 4=completely disabled, cannot carry on any self care, totally confined to bed/chair; 5=dead. Percentage of participants who had at least one point improvement from baseline at any assessment visit as well as at last study visit was reported. Analysis was performed on the safety population.

End point type	Secondary
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End point timeframe:

Baseline, Day 1 of each 28 day cycle up to end of treatment (up to 46 months)

<b>End point values</b>	Vemurafenib			
Subject group type	Reporting group			
Number of subjects analysed	3219			
Units: Percentage of Participants				
number (not applicable)				
At Least 1 Point Improvement at Any Visit	24.7			
At Least 1 Point Improvement at Last Visit	9.9			

### Statistical analyses

No statistical analyses for this end point

### Secondary: Percentage of Participants who Received any Concomitant Medications

End point title	Percentage of Participants who Received any Concomitant Medications
End point description:	
Concomitant medications were all medications taken during the study, including those started before but ongoing at first dose. No medications for Melanoma were included. Percentage of participants who received at least one concomitant medication was reported. Analysis was performed on the safety population.	
End point type	Secondary
End point timeframe:	
Baseline up to 46 months	

<b>End point values</b>	Vemurafenib			
Subject group type	Reporting group			
Number of subjects analysed	3219			
Units: Percentage of Participants				
number (not applicable)	93.8			

### Statistical analyses

No statistical analyses for this end point

### Secondary: Percentage of Participants With Best Overall Response (BOR) of Confirmed Complete Response or Partial Response, as per Response Evaluation Criteria in Solid Tumors Version 1.1 (RECIST v1.1)

End point title	Percentage of Participants With Best Overall Response (BOR) of Confirmed Complete Response or Partial Response, as per Response Evaluation Criteria in Solid Tumors Version 1.1 (RECIST v1.1)
End point description:	
BOR was assessed by the investigator according to RECIST v1.1. BOR was defined as having confirmed CR or PR. CR: disappearance of all target and non-target lesions and no new lesions, all pathological lymph nodes must have decreased to less than (<) 10 millimeter (mm) in short axis; PR: at least a 30%	

decrease in the sum of diameters of target lesions (taking as reference the baseline sum diameters), no progression in non-target lesion, and no new lesions. Confirmed responses were those that persisted on repeat imaging greater than or equal to ( $\geq$ ) 4 weeks after initial response. Analysis was performed on the safety population. Number of subjects analyzed = participants with measurable disease at baseline

End point type	Secondary
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End point timeframe:

Baseline until first documentation of confirmed CR or PR (assessed at baseline, at Weeks 8, 16, as per institution standard of care thereafter but a minimum every 16 weeks thereafter until the end of the study [up to 46 months])

<b>End point values</b>	Vemurafenib			
Subject group type	Reporting group			
Number of subjects analysed	2982			
Units: Percentage of Participants				
number (confidence interval 95%)	33.4 (31.7 to 35.1)			

## Statistical analyses

No statistical analyses for this end point

## Secondary: Duration of Response

End point title	Duration of Response
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End point description:

The duration of response was defined as the time between the date of first confirmed CR or PR and date of first progression of disease (PD), or death, from any cause. Responses were assessed as per RECIST v1.1. CR: disappearance of all target and non-target lesions and no new lesions, all pathological lymph nodes must have decreased to  $< 10$  mm in short axis; PR: at least a 30% decrease in the sum of diameters of target lesions (taking as reference the baseline sum diameters), no progression in non-target lesion, and no new lesions. Confirmed responses were those that persisted on repeat imaging  $\geq 4$  weeks after initial response. PD: at least 20% increase in the sum of diameters of target lesions compared to smallest sum of diameters on study and absolute increase of at least 5 mm, progression of existing non target lesions, or presence of new lesion. Analysis was performed on the safety population. Number of subjects analyzed=participants who achieved CR or PR.

End point type	Secondary
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End point timeframe:

From 1st documentation of confirmed CR or PR to PD or death, whichever occurred first (assessed at baseline, at Weeks 8, 16, as per institution standard of care thereafter but a minimum every 16 weeks thereafter until end of the study [up to 46 months])

<b>End point values</b>	Vemurafenib			
Subject group type	Reporting group			
Number of subjects analysed	996			
Units: Months				
median (confidence interval 95%)	7.4 (7.1 to 8.3)			

## Statistical analyses

No statistical analyses for this end point

### Secondary: Time to Response

End point title	Time to Response
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End point description:

Time to response was defined as the time between the date of first treatment and date of first confirmed CR or PR (assessed as per RECIST v1.1). CR: disappearance of all target and non-target lesions and no new lesions, all pathological lymph nodes must have decreased to < 10 mm in short axis; PR: at least a 30% decrease in the sum of diameters of target lesions (taking as reference the baseline sum diameters), no progression in non-target lesion, and no new lesions. Confirmed responses were those that persisted on repeat imaging  $\geq$  4 weeks after initial response. Analysis was performed on the safety population. Number of subjects analyzed = participants with measurable disease at baseline.

End point type	Secondary
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End point timeframe:

Baseline until first documentation of confirmed CR or PR, whichever occurred first (assessed at baseline, at Weeks 8, 16, as per institution standard of care thereafter but a minimum every 16 weeks thereafter until the end of the study [up to 46 months])

<b>End point values</b>	Vemurafenib			
Subject group type	Reporting group			
Number of subjects analysed	2982			
Units: Months				
median (full range (min-max))	1.84 (0.8 to 26.7)			

## Statistical analyses

No statistical analyses for this end point

### Secondary: Percentage of Participants With PD Assessed According to RECIST v1.1 or Death

End point title	Percentage of Participants With PD Assessed According to RECIST v1.1 or Death
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End point description:

PD was assessed according to RECIST v1.1. PD was defined as at least 20% increase in the sum of diameters of target lesions compared to smallest sum of diameters on study and absolute increase of at least 5 mm, progression of existing non-target lesions, or presence of new lesion. Analysis was performed on the safety population.

End point type	Secondary
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End point timeframe:

Baseline until PD or death, whichever occurred first (assessed at baseline, at Weeks 8, 16, as per institution standard of care thereafter but a minimum every 16 weeks thereafter until the end of the study [up to 46 months])

<b>End point values</b>	Vemurafenib			
Subject group type	Reporting group			
Number of subjects analysed	3219			
Units: Percentage of Participants				
number (not applicable)	87.3			

### Statistical analyses

No statistical analyses for this end point

### Secondary: Progression Free Survival (PFS)

End point title | Progression Free Survival (PFS)

End point description:

PFS was defined as the time between the date of the first treatment and the date of first progression or death from any cause. PD was assessed according to RECIST v1.1. PD was defined as at least 20% increase in the sum of diameters of target lesions compared to smallest sum of diameters on study and absolute increase of at least 5 mm, progression of existing non-target lesions, or presence of new lesion. Analysis was performed on the safety population.

End point type | Secondary

End point timeframe:

Baseline until PD or death, whichever occurred first (assessed at baseline, at Weeks 8, 16, as per institution standard of care thereafter but a minimum every 16 weeks thereafter until the end of the study [up to 46 months])

<b>End point values</b>	Vemurafenib			
Subject group type	Reporting group			
Number of subjects analysed	3219			
Units: Months				
median (confidence interval 95%)	5.6 (5.5 to 5.8)			

### Statistical analyses

No statistical analyses for this end point

### Secondary: Percentage of Participants Who Died

End point title | Percentage of Participants Who Died

End point description:

Analysis was performed on the safety population.

End point type	Secondary
End point timeframe:	
Baseline until death (maximum up to 46 months)	

<b>End point values</b>	Vemurafenib			
Subject group type	Reporting group			
Number of subjects analysed	3219			
Units: Percentage of participants				
number (not applicable)	63.8			

### Statistical analyses

No statistical analyses for this end point

### Secondary: Overall Survival (OS)

End point title	Overall Survival (OS)
End point description:	
Overall Survival was defined as the time from the date of first treatment to the date of death, regardless of the cause of death. Analysis was performed on the safety population.	
End point type	Secondary
End point timeframe:	
Baseline until death (maximum up to 46 months)	

<b>End point values</b>	Vemurafenib			
Subject group type	Reporting group			
Number of subjects analysed	3219			
Units: Months				
median (confidence interval 95%)	12.1 (11.5 to 12.7)			

### Statistical analyses

No statistical analyses for this end point

## Adverse events

### Adverse events information

Timeframe for reporting adverse events:

Baseline up to 28 days post end of treatment (maximum up to 46 months)

Assessment type	Non-systematic
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### Dictionary used

Dictionary name	MedDRA
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Dictionary version	18.1
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### Reporting groups

Reporting group title	Vemurafenib
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Reporting group description:

Participants received continuous oral doses of vemurafenib 960 mg (four 240 mg tablets) twice daily in each 28-day treatment cycle until the development of progressive disease, unacceptable toxicity, consent withdrawal, protocol violations endangering participant's safety, death or study termination by the Sponsor.

Serious adverse events	Vemurafenib		
Total subjects affected by serious adverse events			
subjects affected / exposed	1114 / 3219 (34.61%)		
number of deaths (all causes)	2054		
number of deaths resulting from adverse events			
Neoplasms benign, malignant and unspecified (incl cysts and polyps)			
Basal cell carcinoma			
subjects affected / exposed	68 / 3219 (2.11%)		
occurrences causally related to treatment / all	68 / 79		
deaths causally related to treatment / all	0 / 0		
Chronic lymphocytic leukaemia			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Intracranial tumour haemorrhage			
subjects affected / exposed	7 / 3219 (0.22%)		
occurrences causally related to treatment / all	4 / 7		
deaths causally related to treatment / all	2 / 3		
Fibrous histiocytoma			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Keratoacanthoma</b>			
subjects affected / exposed	260 / 3219 (8.08%)		
occurrences causally related to treatment / all	359 / 361		
deaths causally related to treatment / all	0 / 0		
<b>Malignant melanoma</b>			
subjects affected / exposed	43 / 3219 (1.34%)		
occurrences causally related to treatment / all	36 / 46		
deaths causally related to treatment / all	0 / 0		
<b>Malignant melanoma in situ</b>			
subjects affected / exposed	10 / 3219 (0.31%)		
occurrences causally related to treatment / all	8 / 10		
deaths causally related to treatment / all	0 / 0		
<b>Melanocytic naevus</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	2 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Mycosis Fungoides</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	2 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Papilloma</b>			
subjects affected / exposed	4 / 3219 (0.12%)		
occurrences causally related to treatment / all	5 / 5		
deaths causally related to treatment / all	0 / 0		
<b>Skin papilloma</b>			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	3 / 3		
deaths causally related to treatment / all	0 / 0		
<b>Squamous cell carcinoma of skin</b>			

subjects affected / exposed	259 / 3219 (8.05%)		
occurrences causally related to treatment / all	348 / 352		
deaths causally related to treatment / all	0 / 0		
Tumour haemorrhage			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	1 / 1		
Tumour Ulceration			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Tumour necrosis			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Acanthoma			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	3 / 3		
deaths causally related to treatment / all	0 / 0		
Adrenocarcinoma of Colon			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	0 / 0		
Benign Neoplasm of Skin			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Bowen's Disease			
subjects affected / exposed	5 / 3219 (0.16%)		
occurrences causally related to treatment / all	7 / 7		
deaths causally related to treatment / all	0 / 0		
Breast Cancer			

subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Carcinoma In Situ of Skin</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Cervix Carcinoma Stage 0</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Cholesteatoma</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Chondrosarcoma</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
<b>Ependymoma</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Eye Naevus</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Infected Neoplasm</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Intraductal Proliferative Breast Lesion</b>			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Invasive Ductal Breast Carcinoma</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Kaposi's Sarcoma</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Lymphoma</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Myxoid Liposarcoma</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
<b>Neoplasm</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Neuroendocrine Carcinoma</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Neuroendocrine Carcinoma of the Skin</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Neuroendocrine Tumour</b>			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Oral Papilloma</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Pancreatic Carcinoma</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	2 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Rectal Cancer</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Seborrhoeic Keratosis</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Skin Neoplasm Bleeding</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Squamous Cell Carcinoma of the Oral Cavity</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Squamous Cell Carcinoma of the Vulva</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Superficial Spreading Melanoma Stage Unspecified</b>			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Synovial Sarcoma</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Transitional Cell Carcinoma</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	2 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Tumour Associated Fever</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Lentigo Maligna</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Vascular disorders</b>			
<b>Intermittent Claudication</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Extremity necrosis</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
<b>Hypotension</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
thrombophlebitis			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Circulatory Collapse</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
<b>Deep vein thrombosis</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Haematoma</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Hypertension</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Hypertensive Crisis</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
<b>General disorders and administration site conditions</b>			
<b>Chest pain</b>			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 0		
<b>Chills</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Chronic fatigue syndrome</b>			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	1 / 1		
<b>Death</b>			
subjects affected / exposed	7 / 3219 (0.22%)		
occurrences causally related to treatment / all	1 / 7		
deaths causally related to treatment / all	1 / 7		
<b>Drug Dislocation</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Drug interaction</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Fatigue</b>			
subjects affected / exposed	6 / 3219 (0.19%)		
occurrences causally related to treatment / all	5 / 8		
deaths causally related to treatment / all	0 / 0		
<b>General physical health deterioration</b>			
subjects affected / exposed	9 / 3219 (0.28%)		
occurrences causally related to treatment / all	4 / 9		
deaths causally related to treatment / all	2 / 5		
<b>Malaise</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	2 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Multi-organ disorder</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	1 / 1		
<b>Multi-organ failure</b>			

subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	2 / 2		
deaths causally related to treatment / all	2 / 2		
<b>Oedema</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Oedema peripheral</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Pyrexia</b>			
subjects affected / exposed	32 / 3219 (0.99%)		
occurrences causally related to treatment / all	17 / 34		
deaths causally related to treatment / all	0 / 0		
<b>Sudden cardiac death</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
<b>Asthenia</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Generalised Oedema</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Impaired Healing</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Surgical Failure</b>			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Visceral Pain			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Euthanasia			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 2		
Immune system disorders			
Drug hypersensitivity			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Hypersensitivity			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
Food allergy			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Reproductive system and breast disorders			
Cervical polyp			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Uterine Haemorrhage			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		

Adnexal Torsion			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Endometrial Hyperplasia			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Metrorrhagia			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Peyronie's Disease			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Uterine cervical erosion			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Respiratory, thoracic and mediastinal disorders			
Acute respiratory distress syndrome			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
Chronic obstructive pulmonary disease			
subjects affected / exposed	4 / 3219 (0.12%)		
occurrences causally related to treatment / all	1 / 4		
deaths causally related to treatment / all	0 / 1		
Acute respiratory failure			
subjects affected / exposed	4 / 3219 (0.12%)		
occurrences causally related to treatment / all	0 / 4		
deaths causally related to treatment / all	0 / 1		

Dyspnoea				
subjects affected / exposed	4 / 3219 (0.12%)			
occurrences causally related to treatment / all	1 / 4			
deaths causally related to treatment / all	1 / 1			
Epistaxis				
subjects affected / exposed	1 / 3219 (0.03%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Haemoptysis				
subjects affected / exposed	2 / 3219 (0.06%)			
occurrences causally related to treatment / all	1 / 2			
deaths causally related to treatment / all	0 / 0			
Pleural effusion				
subjects affected / exposed	6 / 3219 (0.19%)			
occurrences causally related to treatment / all	2 / 6			
deaths causally related to treatment / all	0 / 0			
Pneumothorax				
subjects affected / exposed	1 / 3219 (0.03%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Pulmonary embolism				
subjects affected / exposed	13 / 3219 (0.40%)			
occurrences causally related to treatment / all	3 / 13			
deaths causally related to treatment / all	0 / 3			
Respiratory failure				
subjects affected / exposed	1 / 3219 (0.03%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Acute Pulmonary Oedema				
subjects affected / exposed	2 / 3219 (0.06%)			
occurrences causally related to treatment / all	0 / 2			
deaths causally related to treatment / all	0 / 1			
Bronchial Haemorrhage				

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
hydrothorax			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Pleurisy			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Pleuritic Pain			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Pneumonitis			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Pulmonary Toxicity			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	2 / 2		
deaths causally related to treatment / all	1 / 1		
Vocal Cord Leukoplakia			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Psychiatric disorders			
Anxiety			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 0		
Completed suicide			

subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 2		
<b>Confusional state</b>			
subjects affected / exposed	5 / 3219 (0.16%)		
occurrences causally related to treatment / all	0 / 5		
deaths causally related to treatment / all	0 / 0		
<b>Psychotic disorder</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Suicidal ideation</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Bipolar Disorder</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Delirium</b>			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 0		
<b>Mania</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Mood Altered</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Investigations</b>			
Aspartate aminotransferase increased			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Blood creatinine increased			
subjects affected / exposed	7 / 3219 (0.22%)		
occurrences causally related to treatment / all	3 / 7		
deaths causally related to treatment / all	0 / 0		
Gamma glutamyltransferase increased			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	2 / 2		
deaths causally related to treatment / all	0 / 0		
C-reactive protein increased			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Hepatic enzyme increased			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Lipase increased			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Liver function test abnormal			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	3 / 3		
deaths causally related to treatment / all	0 / 0		
Blood glucose increased			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
Electrocardiogram QT prolonged			

subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	4 / 4		
deaths causally related to treatment / all	0 / 0		
<b>Injury, poisoning and procedural complications</b>			
Alcohol Poisoning			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
Fall			
subjects affected / exposed	4 / 3219 (0.12%)		
occurrences causally related to treatment / all	1 / 4		
deaths causally related to treatment / all	0 / 0		
Femoral neck fracture			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 0		
Over dose			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	0 / 0		
Post procedural haemorrhage			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
Hip fracture			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Radiation injury			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
Respiratory fume inhalation disorder			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Spinal fracture</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Cystitis Radiation</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Facial Bone Fracture</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Femur Fracture</b>			
subjects affected / exposed	4 / 3219 (0.12%)		
occurrences causally related to treatment / all	0 / 4		
deaths causally related to treatment / all	0 / 0		
<b>Fracture</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Fracture Displacement</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Head Injury</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
<b>Humerus Fracture</b>			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Lumbar Vertrebral Fracture</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Multiple Injuries</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
<b>Patella Fracture</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Radiation Necrosis</b>			
subjects affected / exposed	4 / 3219 (0.12%)		
occurrences causally related to treatment / all	0 / 4		
deaths causally related to treatment / all	0 / 0		
<b>Radius Fracture</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Road Traffic Accident</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Spinal Compression Fracture</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Splenic Rupture</b>			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Subdural Haematoma</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Sunburn</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Tendon Injury</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Tibia Fracture</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Toxicity to Various Agents</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	2 / 3		
deaths causally related to treatment / all	0 / 0		
<b>Cardiac disorders</b>			
<b>Acute myocardial infarction</b>			
subjects affected / exposed	10 / 3219 (0.31%)		
occurrences causally related to treatment / all	1 / 10		
deaths causally related to treatment / all	1 / 4		
<b>Angina pectoris</b>			
subjects affected / exposed	5 / 3219 (0.16%)		
occurrences causally related to treatment / all	0 / 5		
deaths causally related to treatment / all	0 / 0		
<b>Arrhythmia</b>			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Atrial fibrillation</b>			
subjects affected / exposed	15 / 3219 (0.47%)		
occurrences causally related to treatment / all	6 / 15		
deaths causally related to treatment / all	0 / 0		
<b>Atrial flutter</b>			
subjects affected / exposed	4 / 3219 (0.12%)		
occurrences causally related to treatment / all	2 / 4		
deaths causally related to treatment / all	0 / 0		
<b>Cardiac failure</b>			
subjects affected / exposed	6 / 3219 (0.19%)		
occurrences causally related to treatment / all	3 / 6		
deaths causally related to treatment / all	0 / 0		
<b>Coronary artery disease</b>			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	1 / 3		
deaths causally related to treatment / all	0 / 1		
<b>Left ventricular dysfunction</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Myocardial infarction</b>			
subjects affected / exposed	7 / 3219 (0.22%)		
occurrences causally related to treatment / all	1 / 8		
deaths causally related to treatment / all	0 / 2		
<b>Pericardial effusion</b>			
subjects affected / exposed	7 / 3219 (0.22%)		
occurrences causally related to treatment / all	4 / 7		
deaths causally related to treatment / all	0 / 0		
<b>Pericardial haemorrhage</b>			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Pericarditis</b>			
subjects affected / exposed	7 / 3219 (0.22%)		
occurrences causally related to treatment / all	7 / 8		
deaths causally related to treatment / all	0 / 0		
<b>Tachycardia</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Torsade de pointes</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	1 / 1		
<b>Acute Coronary Syndrome</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Atrioventricular Block</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Angina Unstable</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Cardiac Arrest</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 2		
<b>Cardiac Failure Acute</b>			

subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Cardiac Failure Congestive</b>			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 0		
<b>Cardiogenic Shock</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	1 / 1		
<b>Cardiomyopathy</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Coronary Artery Stenosis</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Sinus Node Dysfunction</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Supraventricular Tachycardia</b>			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	1 / 4		
deaths causally related to treatment / all	0 / 0		
<b>Nervous system disorders</b>			
<b>Brain oedema</b>			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	0 / 4		
deaths causally related to treatment / all	0 / 0		
<b>Cerebral haematoma</b>			

subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	2 / 3		
deaths causally related to treatment / all	1 / 1		
<b>Cerebral haemorrhage</b>			
subjects affected / exposed	14 / 3219 (0.43%)		
occurrences causally related to treatment / all	4 / 14		
deaths causally related to treatment / all	3 / 8		
<b>Cognitive disorder</b>			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	1 / 3		
deaths causally related to treatment / all	0 / 0		
<b>Cerebrovascular accident</b>			
subjects affected / exposed	10 / 3219 (0.31%)		
occurrences causally related to treatment / all	1 / 10		
deaths causally related to treatment / all	0 / 5		
<b>Depressed Level of Consciousness</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
<b>Dizziness</b>			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	2 / 3		
deaths causally related to treatment / all	0 / 0		
<b>Dysarthria</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Epilepsy</b>			
subjects affected / exposed	10 / 3219 (0.31%)		
occurrences causally related to treatment / all	1 / 14		
deaths causally related to treatment / all	0 / 0		
<b>Generalised Tonic Clonic Seizure</b>			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Facial paresis			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Haemorrhage intracranial			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 2		
Haemorrhagic stroke			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	1 / 1		
Headache			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	2 / 3		
deaths causally related to treatment / all	0 / 0		
Intracranial Pressure Increased			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Hydrocephalus			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 1		
Ischaemic stroke			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
Nervous system disorder			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Neuralgia</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	2 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Neuropathy peripheral</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Sciatica</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Stupor</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Syncope</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Transient ischaemic attack</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Tremor</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>VIIth nerve paralysis</b>			

subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Aphasia</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Balance Disorder</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Carpal Tunnel Syndrome</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Cerebral Disorder</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Cerebrospinal Fluid Leakage</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Facial Nerve Disorder</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Metabolic Encephalopathy</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Migraine</b>			

subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Monoplegia</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Neurotoxicity</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Partial Seizures</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Peripheral Sensorimotor Neuropathy</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	2 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Polyneuropathy</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Seizure</b>			
subjects affected / exposed	12 / 3219 (0.37%)		
occurrences causally related to treatment / all	0 / 13		
deaths causally related to treatment / all	0 / 0		
<b>Simple Partial Seizure</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Spinal Cord Compression</b>			

subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 0		
<b>Status Epilepticus</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Subarachnoid Haemorrhage</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Vasogenic Cerebral Oedema</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Cerebellar Infarction</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Blood and lymphatic system disorders</b>			
<b>Anaemia</b>			
subjects affected / exposed	22 / 3219 (0.68%)		
occurrences causally related to treatment / all	7 / 25		
deaths causally related to treatment / all	0 / 0		
<b>Disseminated intravascular coagulation</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	1 / 2		
<b>Eosinophilia</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Neutropenia</b>			

subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	2 / 3		
deaths causally related to treatment / all	0 / 0		
<b>Thrombocytopenia</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Agranulocytosis</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
<b>Febrile Neutropenia</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Lymph Node Pain</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Microcytic Anaemia</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Ear and labyrinth disorders</b>			
<b>Deafness</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Vertigo</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Hearnig Impaired</b>			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Eye disorders</b>			
<b>Neovascular Age-related macular degeneration</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Diplopia</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Iridocyclitis</b>			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	3 / 3		
deaths causally related to treatment / all	0 / 0		
<b>Uveitis</b>			
subjects affected / exposed	6 / 3219 (0.19%)		
occurrences causally related to treatment / all	6 / 6		
deaths causally related to treatment / all	0 / 0		
<b>Vitreous haemorrhage</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Cataract</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	2 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Glaucoma</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Macular Oedema</b>			

subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Papilloedema</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Retinal Artery occlusion</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Retinal Detachment</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Retinopathy</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Iritis</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Gastrointestinal disorders</b>			
<b>Abdominal distension</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Abdominal pain</b>			
subjects affected / exposed	4 / 3219 (0.12%)		
occurrences causally related to treatment / all	1 / 4		
deaths causally related to treatment / all	0 / 0		
<b>Abdominal pain lower</b>			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Abdominal pain upper			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Ascites			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Constipation			
subjects affected / exposed	6 / 3219 (0.19%)		
occurrences causally related to treatment / all	1 / 6		
deaths causally related to treatment / all	0 / 0		
Diarrhoea			
subjects affected / exposed	16 / 3219 (0.50%)		
occurrences causally related to treatment / all	9 / 16		
deaths causally related to treatment / all	0 / 0		
Diverticular perforation			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
Duodenal ulcer			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
Gastrointestinal haemorrhage			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 0		
Gastrooesophageal reflux disease			

subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Heamatemesis</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Ileus</b>			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 0		
<b>Inguinal hernia</b>			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 0		
<b>Intestinal obstruction</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Intestinal perforation</b>			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	1 / 3		
deaths causally related to treatment / all	0 / 0		
<b>Jejunal perforation</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Nausea</b>			
subjects affected / exposed	13 / 3219 (0.40%)		
occurrences causally related to treatment / all	11 / 15		
deaths causally related to treatment / all	0 / 0		
<b>Oesophagitis</b>			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Pancreatitis</b>			
subjects affected / exposed	4 / 3219 (0.12%)		
occurrences causally related to treatment / all	2 / 4		
deaths causally related to treatment / all	0 / 0		
<b>Pancreatitis acute</b>			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	1 / 3		
deaths causally related to treatment / all	0 / 0		
<b>Small intestinal obstruction</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Vomiting</b>			
subjects affected / exposed	17 / 3219 (0.53%)		
occurrences causally related to treatment / all	9 / 18		
deaths causally related to treatment / all	0 / 0		
<b>Anal Fistula</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Anal skin Tags</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Colitis</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Dyspepsia</b>			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Dysphagia</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Enterocolitis</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Faeces Pale</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Food Poisoning</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Gastric Haemorrhagic</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Gastrointestinal Necrosis</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
<b>Gastrointestinal Toxicity</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Haemorrhoidal haemorrhage</b>			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Intestinal Ischaemia</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
<b>Intussusception</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Large intestine perforation</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Leukoplakia oral</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	2 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Lower gastrointestinal haemorrhage</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Melaena</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Mouth Ulceration</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Paraesthesia oral</b>			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Rectal Haemorrhage			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Retroperitoneal haematoma			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Hepatobiliary disorders			
Cholecystitis acute			
subjects affected / exposed	4 / 3219 (0.12%)		
occurrences causally related to treatment / all	0 / 4		
deaths causally related to treatment / all	0 / 0		
Cholelithiasis			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Drug-induced liver injury			
subjects affected / exposed	4 / 3219 (0.12%)		
occurrences causally related to treatment / all	5 / 5		
deaths causally related to treatment / all	0 / 0		
Hepatic function abnormal			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Hepatitis			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Hepatotoxicity			

subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	3 / 3		
deaths causally related to treatment / all	0 / 0		
<b>Cholangitis</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Cholecystitis</b>			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	1 / 3		
deaths causally related to treatment / all	0 / 0		
<b>Hepatic Failure</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Hepatitis Toxic</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Skin and subcutaneous tissue disorders</b>			
<b>Acanthosis</b>			
subjects affected / exposed	4 / 3219 (0.12%)		
occurrences causally related to treatment / all	6 / 6		
deaths causally related to treatment / all	0 / 0		
<b>Actinic keratosis</b>			
subjects affected / exposed	9 / 3219 (0.28%)		
occurrences causally related to treatment / all	10 / 10		
deaths causally related to treatment / all	0 / 0		
<b>Angioedema</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	2 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Dermatitis exfoliative</b>			

subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	4 / 4		
deaths causally related to treatment / all	0 / 0		
Hypersensitivity Vasculitis			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Photosensitivity reaction			
subjects affected / exposed	5 / 3219 (0.16%)		
occurrences causally related to treatment / all	5 / 5		
deaths causally related to treatment / all	0 / 0		
Rash maculo-papular			
subjects affected / exposed	7 / 3219 (0.22%)		
occurrences causally related to treatment / all	7 / 7		
deaths causally related to treatment / all	0 / 0		
Rash			
subjects affected / exposed	5 / 3219 (0.16%)		
occurrences causally related to treatment / all	4 / 5		
deaths causally related to treatment / all	0 / 0		
Rash pruritic			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Skin Mass			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Toxic epidermal necrolysis			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	3 / 3		
deaths causally related to treatment / all	1 / 1		
Lichenoid Keratosis			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Dermal Cyst</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Dermatitis exfoliative Generalised</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Drug eruption</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	2 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Drug reaction with eosinophilia and systemic symptoms</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	2 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Erythema Nodosum</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Hyperkeratosis</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Panniculitis</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	2 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Rash Erythematous</b>			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Rash Follicular</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Rash Generalised</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	2 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Rash Macular</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Rash Papular</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Steven Johnson Syndrome</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Renal and urinary disorders</b>			
<b>Calculus ureteric</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Glomerulonephritis minimal lesion</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Renal Colic</b>			

subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Haematuria</b>			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 0		
<b>Urinary tract disorder</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Acute Kidney Injury</b>			
subjects affected / exposed	8 / 3219 (0.25%)		
occurrences causally related to treatment / all	4 / 9		
deaths causally related to treatment / all	0 / 0		
<b>Nephrolithiasis</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 0		
<b>Nephropathy</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Nephropathy Toxic</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Tubulointerstitial Nephritis</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Urinary Fistula</b>			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Endocrine disorders</b>			
<b>Hyperadrenalism</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Musculoskeletal and connective tissue disorders</b>			
<b>Arthralgia</b>			
subjects affected / exposed	12 / 3219 (0.37%)		
occurrences causally related to treatment / all	11 / 13		
deaths causally related to treatment / all	0 / 0		
<b>Back pain</b>			
subjects affected / exposed	6 / 3219 (0.19%)		
occurrences causally related to treatment / all	1 / 6		
deaths causally related to treatment / all	0 / 0		
<b>Bursitis</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Muscular weakness</b>			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	1 / 3		
deaths causally related to treatment / all	0 / 0		
<b>Musculoskeletal chest pain</b>			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	0 / 5		
deaths causally related to treatment / all	0 / 0		
<b>Arthritis</b>			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	2 / 3		
deaths causally related to treatment / all	0 / 0		

Myalgia				
subjects affected / exposed	2 / 3219 (0.06%)			
occurrences causally related to treatment / all	1 / 2			
deaths causally related to treatment / all	0 / 0			
Neck pain				
subjects affected / exposed	1 / 3219 (0.03%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Osteoporosis				
subjects affected / exposed	1 / 3219 (0.03%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Pathological fracture				
subjects affected / exposed	1 / 3219 (0.03%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Dupuytren's Contracture				
subjects affected / exposed	1 / 3219 (0.03%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Osteitis				
subjects affected / exposed	1 / 3219 (0.03%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Pain in extremity				
subjects affected / exposed	1 / 3219 (0.03%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Spinal Pain				
subjects affected / exposed	1 / 3219 (0.03%)			
occurrences causally related to treatment / all	0 / 2			
deaths causally related to treatment / all	0 / 0			
Osteonecrosis				

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Infections and infestations</b>			
<b>Abscess Limb</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Anal abscess</b>			
subjects affected / exposed	5 / 3219 (0.16%)		
occurrences causally related to treatment / all	2 / 6		
deaths causally related to treatment / all	0 / 0		
<b>Blastocystis infection</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Bronchitis</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Cellulitis</b>			
subjects affected / exposed	5 / 3219 (0.16%)		
occurrences causally related to treatment / all	2 / 6		
deaths causally related to treatment / all	0 / 0		
<b>Clostridium difficile infection</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Device related infection</b>			
subjects affected / exposed	4 / 3219 (0.12%)		
occurrences causally related to treatment / all	0 / 4		
deaths causally related to treatment / all	0 / 0		
<b>Diverticulitis</b>			

subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 0		
<b>Erysipelas</b>			
subjects affected / exposed	9 / 3219 (0.28%)		
occurrences causally related to treatment / all	1 / 10		
deaths causally related to treatment / all	0 / 0		
<b>Escherichia bacteraemia</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Escherichia sepsis</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 1		
<b>Gastroenteritis</b>			
subjects affected / exposed	4 / 3219 (0.12%)		
occurrences causally related to treatment / all	1 / 4		
deaths causally related to treatment / all	0 / 0		
<b>Groin abscess</b>			
subjects affected / exposed	5 / 3219 (0.16%)		
occurrences causally related to treatment / all	1 / 5		
deaths causally related to treatment / all	0 / 0		
<b>Herpes zoster</b>			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 0		
<b>H1N1 Influenza</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Infection</b>			

subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	0 / 0		
Lower respiratory tract infection			
subjects affected / exposed	7 / 3219 (0.22%)		
occurrences causally related to treatment / all	0 / 7		
deaths causally related to treatment / all	0 / 0		
Lung infection			
subjects affected / exposed	6 / 3219 (0.19%)		
occurrences causally related to treatment / all	0 / 6		
deaths causally related to treatment / all	0 / 0		
Oesophageal candidiasis			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 0		
Oral candidiasis			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Otitis externa			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Pilonidal cyst			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	0 / 0		
Pharyngotonsillitis			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Pneumonia			

subjects affected / exposed	31 / 3219 (0.96%)		
occurrences causally related to treatment / all	4 / 39		
deaths causally related to treatment / all	1 / 3		
Pneumocystis jiroveci pneumonia			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 1		
Postoperative Wound Infection			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
Rash pustular			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Respiratory tract infection			
subjects affected / exposed	7 / 3219 (0.22%)		
occurrences causally related to treatment / all	0 / 7		
deaths causally related to treatment / all	0 / 2		
Sepsis			
subjects affected / exposed	7 / 3219 (0.22%)		
occurrences causally related to treatment / all	1 / 7		
deaths causally related to treatment / all	1 / 3		
Salmonella sepsis			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Septic shock			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Skin infection			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Subcutaneous abscess</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Upper respiratory tract infection</b>			
subjects affected / exposed	4 / 3219 (0.12%)		
occurrences causally related to treatment / all	0 / 4		
deaths causally related to treatment / all	0 / 0		
<b>Urinary tract infection</b>			
subjects affected / exposed	11 / 3219 (0.34%)		
occurrences causally related to treatment / all	0 / 11		
deaths causally related to treatment / all	0 / 0		
<b>Urosepsis</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Viral infection</b>			
subjects affected / exposed	3 / 3219 (0.09%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 1		
<b>Vulvitis</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Wound infection</b>			
subjects affected / exposed	4 / 3219 (0.12%)		
occurrences causally related to treatment / all	0 / 4		
deaths causally related to treatment / all	0 / 0		
<b>Wound infection staphylococcal</b>			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Acinetobacter Infection</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Appendicitis</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Appendicitis Perforated</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Arthritis Bacterial</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Brain Abscess</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Atypical Pneumonia</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Cholecystitis Infective</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Colonic Abscess</b>			

subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 0		
<b>Cystitis</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Gastroenteritis Viral</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Gastrointestinal Infection</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Hypopyon</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Infected Dermal Cyst</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Infectious Pleural Effusion</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Localised Infection</b>			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
<b>Lymph Gland Infection</b>			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Ophthalmic Herpes Zoster</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Perineal Abscess</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Skin Bacterial Infection</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Soft Tissue Infection</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
<b>Subdiaphragmatic Abscess</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 0		
<b>Tonsillitis</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Tooth Abscess</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Tooth Infection</b>			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Clostridium Colitis			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Metabolism and nutrition disorders			
Cachexia			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Dehydration			
subjects affected / exposed	4 / 3219 (0.12%)		
occurrences causally related to treatment / all	1 / 4		
deaths causally related to treatment / all	0 / 0		
Diabetes mellitus			
subjects affected / exposed	4 / 3219 (0.12%)		
occurrences causally related to treatment / all	0 / 4		
deaths causally related to treatment / all	0 / 0		
Diabetic ketoacidosis			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	0 / 0		
Hyperuricaemia			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	0 / 0		
Hyperglycaemia			
subjects affected / exposed	2 / 3219 (0.06%)		
occurrences causally related to treatment / all	2 / 2		
deaths causally related to treatment / all	0 / 0		
Hypokalaemia			

subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Hyponatraemia</b>			
subjects affected / exposed	4 / 3219 (0.12%)		
occurrences causally related to treatment / all	0 / 4		
deaths causally related to treatment / all	0 / 0		
<b>Tumour lysis syndrome</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Decreased Appetite</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Hypoglycaemia</b>			
subjects affected / exposed	4 / 3219 (0.12%)		
occurrences causally related to treatment / all	1 / 4		
deaths causally related to treatment / all	0 / 0		
<b>Type 2 Diabestes Mellitus</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Electrolyte Imbalance</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
<b>Hypertriglyceridaemia</b>			
subjects affected / exposed	1 / 3219 (0.03%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		

Frequency threshold for reporting non-serious adverse events: 5 %

<b>Non-serious adverse events</b>	Vemurafenib		
Total subjects affected by non-serious adverse events			
subjects affected / exposed	2956 / 3219 (91.83%)		
Neoplasms benign, malignant and unspecified (incl cysts and polyps)			
Skin papilloma			
subjects affected / exposed	621 / 3219 (19.29%)		
occurrences (all)	953		
Melanocytic Naevus			
subjects affected / exposed	227 / 3219 (7.05%)		
occurrences (all)	332		
Seborrhoeic Keratosis			
subjects affected / exposed	266 / 3219 (8.26%)		
occurrences (all)	333		
Vascular disorders			
Hypertension			
subjects affected / exposed	264 / 3219 (8.20%)		
occurrences (all)	348		
General disorders and administration site conditions			
Fatigue			
subjects affected / exposed	834 / 3219 (25.91%)		
occurrences (all)	1129		
Pyrexia			
subjects affected / exposed	363 / 3219 (11.28%)		
occurrences (all)	484		
Asthenia			
subjects affected / exposed	382 / 3219 (11.87%)		
occurrences (all)	535		
Oedema Peripheral			
subjects affected / exposed	238 / 3219 (7.39%)		
occurrences (all)	274		
Respiratory, thoracic and mediastinal disorders			

Cough subjects affected / exposed occurrences (all)	191 / 3219 (5.93%) 215		
Psychiatric disorders Insomnia subjects affected / exposed occurrences (all)	163 / 3219 (5.06%) 175		
Investigations Weight decreased subjects affected / exposed occurrences (all)	414 / 3219 (12.86%) 485		
Electrocardiogram QT prolonged subjects affected / exposed occurrences (all)	521 / 3219 (16.19%) 753		
Injury, poisoning and procedural complications Sunburn subjects affected / exposed occurrences (all)	318 / 3219 (9.88%) 522		
Nervous system disorders Headache subjects affected / exposed occurrences (all)	458 / 3219 (14.23%) 637		
Dysgeusia subjects affected / exposed occurrences (all)	191 / 3219 (5.93%) 205		
Blood and lymphatic system disorders Anaemia subjects affected / exposed occurrences (all)	215 / 3219 (6.68%) 316		
Gastrointestinal disorders Diarrhoea subjects affected / exposed occurrences (all)	584 / 3219 (18.14%) 866		
Nausea subjects affected / exposed occurrences (all)	718 / 3219 (22.31%) 965		

Vomiting			
subjects affected / exposed	449 / 3219 (13.95%)		
occurrences (all)	635		
Abdominal Pain			
subjects affected / exposed	170 / 3219 (5.28%)		
occurrences (all)	214		
Constipation			
subjects affected / exposed	196 / 3219 (6.09%)		
occurrences (all)	211		
Skin and subcutaneous tissue disorders			
Alopecia			
subjects affected / exposed	873 / 3219 (27.12%)		
occurrences (all)	942		
Dry skin			
subjects affected / exposed	538 / 3219 (16.71%)		
occurrences (all)	597		
Erythema			
subjects affected / exposed	327 / 3219 (10.16%)		
occurrences (all)	415		
Hyperkeratosis			
subjects affected / exposed	830 / 3219 (25.78%)		
occurrences (all)	1201		
Photosensitivity reaction			
subjects affected / exposed	678 / 3219 (21.06%)		
occurrences (all)	891		
Rash			
subjects affected / exposed	556 / 3219 (17.27%)		
occurrences (all)	709		
Pruritus			
subjects affected / exposed	320 / 3219 (9.94%)		
occurrences (all)	370		
Actinic Keratosis			
subjects affected / exposed	246 / 3219 (7.64%)		
occurrences (all)	405		
Palmar Plantar Erythrodysesthesia			

syndrome			
subjects affected / exposed	193 / 3219 (6.00%)		
occurrences (all)	253		
Rash Erythematous			
subjects affected / exposed	195 / 3219 (6.06%)		
occurrences (all)	246		
Rash Maculo- Papular			
subjects affected / exposed	179 / 3219 (5.56%)		
occurrences (all)	240		
Musculoskeletal and connective tissue disorders			
Arthralgia			
subjects affected / exposed	1357 / 3219 (42.16%)		
occurrences (all)	2178		
Myalgia			
subjects affected / exposed	308 / 3219 (9.57%)		
occurrences (all)	402		
Back Pain			
subjects affected / exposed	191 / 3219 (5.93%)		
occurrences (all)	214		
Musculoskeletal Pain			
subjects affected / exposed	222 / 3219 (6.90%)		
occurrences (all)	277		
Pain in Extremity			
subjects affected / exposed	276 / 3219 (8.57%)		
occurrences (all)	343		
Infections and infestations			
Conjunctivitis			
subjects affected / exposed	170 / 3219 (5.28%)		
occurrences (all)	199		
Nasopharyngitis			
subjects affected / exposed	171 / 3219 (5.31%)		
occurrences (all)	223		
Metabolism and nutrition disorders			
Decreased appetite			
subjects affected / exposed	482 / 3219 (14.97%)		
occurrences (all)	603		



## More information

### Substantial protocol amendments (globally)

Were there any global substantial amendments to the protocol? Yes

Date	Amendment
28 April 2011	<ul style="list-style-type: none"><li>• Based on result of a different study similar participant population was allowed to be enrolled in this study and therefore participants with surgically incurable and unresectable Stage IIIC or Stage IV (AJCC) and participants who had not received prior treatment for metastatic melanoma were allowed into the study (in addition to participants who had received prior treatment).</li><li>• Included participants at least 16 years of age due to the assumed similarities between the biological/physiological and pharmacokinetic/pharmacodynamic characteristics of 16 to 18-year-old participants.</li><li>• Excluded participants with a known hypersensitivity to vemurafenib or another BRAF inhibitor.</li><li>• Excluded participants with congenital long QT syndrome considering the recent data of effect of vemurafenib on the QT interval.</li><li>• Clarified the type of concomitant medication that was prohibited or must be used with caution.</li><li>• Included ECG monitoring to be 28 days after starting treatment with vemurafenib, every 4 weeks for the following 3 months, every 12 weeks thereafter and at the end of study/follow-up visit, following the new safety information.</li><li>• Clarified the definition of the date at which the Screening Period started to be the date at which the first archival tumor tissue sample was sent to the central testing laboratory for BRAF mutation testing, except when a study procedure was performed prior to sending the tumor sample to the testing laboratory, when this date was recorded as the date of the start of Screening.</li><li>• Clarified the reporting of SCC to allow for the proper coding to the preferred term cutaneous SCC and clarified that SCC should be reported as an serious AE only if it meets the definition of an serious AE.</li></ul>
30 November 2011	<ul style="list-style-type: none"><li>• Included a long term follow-up safety phase for 24 months after the last participant enrolled.</li><li>• Added information to the study background.</li><li>• Included tolerability as part of the primary objective of the study, together with safety.</li><li>• Clearly defined the end of treatment visit and the 28 day follow-up visit after discontinuation of vemurafenib and updated the sections referring to visits.</li><li>• Increased the number of participants to be screened and enrolled.</li><li>• Clarified that only a protocol violation that endangered a participant's safety would mandate discontinuation of study treatment.</li><li>• Added the option for participants who developed disease progression but who in the opinion of the investigator would still benefit from continuing vemurafenib treatment, could do so after discussion with the Sponsor.</li><li>• Included that for participants who developed any other suspicious lesions (in addition to SCC), tissue from the lesions was also to be provided for confirmation of diagnosis by a designated central pathology laboratory.</li><li>• Clarified when the study enrolment would end in relation to local regulatory approval and reimbursement of vemurafenib for the treatment of metastatic melanoma.</li><li>• Increased the number of study centers.</li><li>• Clarified the exclusion criteria</li><li>• Clarified the use of limited field radiotherapy to include all palliative reasons if the limited field radiotherapy was not considered a target lesion for the RECIST assessments.</li><li>• Increased the follow-up of chest CT for evaluation of non cutaneous SCC.</li><li>• Clarified that the participants with known or suspected bone metastases should undergo radionuclide bone scan or PET scan at baseline if clinically indicated.</li><li>• Clarified the definition of AE and SAEs.</li><li>• Amended the statistical analyses.</li></ul>

15 August 2012	<ul style="list-style-type: none"> <li>• Added that participant recruitment has been higher than expected with additional countries and sites than originally planned being included. The protocol has been updated to reflect the higher number of participant and revised statistical assumptions. It was estimated that approximately 7400 participants will be screened and approximately 3300 participants with the BRAF V600 mutation will be enrolled in the study and receive treatment with vemurafenib.</li> <li>• Increased the visit window to +/- 5 days to allow flexibility and to be more realistic to clinical routine, and the follow-up visit to after 28 days.</li> <li>• Included an additional head and neck examination for monitoring of non cutaneous SCCs to align with the monitoring guidance of the Zelboraf® (marketed name of vemurafenib) Summary of Product Characteristics, to the Long Term Safety Follow-up Phase, to be performed at 6 months following study drug discontinuation or until initiation of another neo plastic therapy.</li> <li>• Added a dermatological evaluation at 6 months following study drug discontinuation or until initiation of another anti neoplastic therapy for monitoring of cutaneous SCCs to align with the monitoring guidance of the vemurafenib Summary of Product Characteristics</li> <li>• Included for consistency throughout the protocol that as well as cutaneous SCC, BCC and keratoacanthoma and other second primary malignancies were defined as events requiring close monitoring.</li> </ul>
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Notes:

### **Interruptions (globally)**

Were there any global interruptions to the trial? No

### **Limitations and caveats**

None reported