



## Clinical trial results:

**A randomized, double-blinded, regimen controlled, phase II, multicenter study to assess the efficacy and safety of two different vismodegib regimens in patients with multiple basal cell carcinomas**

### Summary

EudraCT number	2012-003305-10
Trial protocol	AT DE NL IT ES FR
Global end of trial date	31 August 2016

### Results information

Result version number	v3 (current)
This version publication date	15 September 2017
First version publication date	13 July 2016
Version creation reason	

### Trial information

#### Trial identification

Sponsor protocol code	MO28295
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#### Additional study identifiers

ISRCTN number	-
ClinicalTrials.gov id (NCT number)	NCT01815840
WHO universal trial number (UTN)	-

Notes:

### Sponsors

Sponsor organisation name	F. Hoffmann-La Roche AG
Sponsor organisation address	Grenzacherstrasse 124, Basel, Switzerland, CH-4070
Public contact	F. Hoffmann-La Roche AG, F. Hoffmann-La Roche AG, +41 616878333, global.trial_information@roche.com
Scientific contact	F. Hoffmann-La Roche AG, F. Hoffmann-La Roche AG, +41 616878333, 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?	No

Notes:

## Results analysis stage

Analysis stage	Final
Date of interim/final analysis	31 August 2016
Is this the analysis of the primary completion data?	No
Global end of trial reached?	Yes
Global end of trial date	31 August 2016
Was the trial ended prematurely?	No

Notes:

## General information about the trial

Main objective of the trial:

This randomized, double-blind, regimen-controlled, phase II, multicenter study assessed the efficacy and safety of two different vismodegib regimens in subjects with multiple basal cell carcinoma. Subjects received vismodegib 150 mg orally once daily either in an intermittent schedule of 12 weeks vismodegib followed by 8 weeks placebo (Arm A) or as 24 weeks induction followed by an intermittent schedule of 8 weeks placebo followed by 8 weeks vismodegib (Arm B). Anticipated time on study treatment was 72 weeks.

Protection of trial subjects:

All study subjects were required to read and sign an informed consent form.

Background therapy: -

Evidence for comparator: -

Actual start date of recruitment	30 April 2013
Long term follow-up planned	No
Independent data monitoring committee (IDMC) involvement?	Yes

Notes:

## Population of trial subjects

### Subjects enrolled per country

Country: Number of subjects enrolled	Netherlands: 10
Country: Number of subjects enrolled	Spain: 23
Country: Number of subjects enrolled	Austria: 12
Country: Number of subjects enrolled	France: 38
Country: Number of subjects enrolled	Germany: 39
Country: Number of subjects enrolled	Italy: 27
Country: Number of subjects enrolled	Canada: 15
Country: Number of subjects enrolled	Mexico: 8
Country: Number of subjects enrolled	Russian Federation: 9
Country: Number of subjects enrolled	United States: 48
Worldwide total number of subjects	229
EEA total number of subjects	149

Notes:

### Subjects enrolled per age group

In utero	0
Preterm newborn - gestational age < 37	0

wk	
Newborns (0-27 days)	0
Infants and toddlers (28 days-23 months)	0
Children (2-11 years)	0
Adolescents (12-17 years)	0
Adults (18-64 years)	127
From 65 to 84 years	94
85 years and over	8

## Subject disposition

### Recruitment

Recruitment details: -

### Pre-assignment

Screening details:

229 subjects were enrolled in 10 countries.

### Period 1

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

### Arms

Are arms mutually exclusive?	Yes
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<b>Arm title</b>	Vismodegib Intermittent Schedule
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Arm description:

Vismodegib intermittent schedule of 12 weeks vismodegib followed by 8 weeks placebo, repeated 3 times with a final course of vismodegib (total 72 weeks), followed by 52 weeks treatment-free follow up

Arm type	Experimental
Investigational medicinal product name	Vismodegib
Investigational medicinal product code	
Other name	Erivedge®
Pharmaceutical forms	Capsule
Routes of administration	Oral use

Dosage and administration details:

Vismodegib 150 mg hard gelatin capsule orally once daily

Investigational medicinal product name	Vismodegib Placebo
Investigational medicinal product code	
Other name	
Pharmaceutical forms	Capsule
Routes of administration	Oral use

Dosage and administration details:

Vismodegib placebo orally once daily

<b>Arm title</b>	Vismodegib Induction Followed by Intermittent Schedule
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Arm description:

Vismodegib beginning with 24 weeks induction followed by intermittent schedule 8 weeks placebo, 8 weeks vismodegib (total 72 weeks), followed by 52 weeks treatment-free follow up

Arm type	Experimental
Investigational medicinal product name	Vismodegib
Investigational medicinal product code	
Other name	Erivedge®
Pharmaceutical forms	Capsule
Routes of administration	Oral use

Dosage and administration details:

Vismodegib 150 mg hard gelatin capsule orally once daily

Investigational medicinal product name	Vismodegib Placebo
Investigational medicinal product code	
Other name	

Pharmaceutical forms	Capsule
Routes of administration	Oral use

Dosage and administration details:

Vismodegib placebo orally once daily

Number of subjects in period 1	Vismodegib Intermittent Schedule	Vismodegib Induction Followed by Intermittent Schedule
Started	116	113
Completed	57	50
Not completed	59	63
Withdrew Consent	31	30
Disease progression	3	2
Sponsor Termination Treatment	1	-
Adverse Event	10	16
Other pre-specified	1	-
Death	2	2
Refused Treatment	5	1
Administrative/Other	1	3
Investigators Decision	2	3
Lost to follow-up	3	6

## Baseline characteristics

### Reporting groups

Reporting group title	Vismodegib Intermittent Schedule
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Reporting group description:

Vismodegib intermittent schedule of 12 weeks vismodegib followed by 8 weeks placebo, repeated 3 times with a final course of vismodegib (total 72 weeks), followed by 52 weeks treatment-free follow up

Reporting group title	Vismodegib Induction Followed by Intermittent Schedule
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Reporting group description:

Vismodegib beginning with 24 weeks induction followed by intermittent schedule 8 weeks placebo, 8 weeks vismodegib (total 72 weeks), followed by 52 weeks treatment-free follow up

Reporting group values	Vismodegib Intermittent Schedule	Vismodegib Induction Followed by Intermittent Schedule	Total
Number of subjects	116	113	229
Age categorical Units: Subjects			

Age continuous Units: years arithmetic mean standard deviation	61.1 ± 13.94	59.9 ± 15.35	-
Gender categorical Units: Subjects			
Female	35	25	60
Male	81	88	169

## End points

### End points reporting groups

Reporting group title	Vismodegib Intermittent Schedule
Reporting group description: Vismodegib intermittent schedule of 12 weeks vismodegib followed by 8 weeks placebo, repeated 3 times with a final course of vismodegib (total 72 weeks), followed by 52 weeks treatment-free follow up	
Reporting group title	Vismodegib Induction Followed by Intermittent Schedule
Reporting group description: Vismodegib beginning with 24 weeks induction followed by intermittent schedule 8 weeks placebo, 8 weeks vismodegib (total 72 weeks), followed by 52 weeks treatment-free follow up	

### Primary: Mean Percent Change From Baseline in the Number of Clinically Evident Basal Cell Carcinomas at Week 73 (After 72 Weeks of Treatment)

End point title	Mean Percent Change From Baseline in the Number of Clinically Evident Basal Cell Carcinomas at Week 73 (After 72 Weeks of Treatment)
End point description: The total number of clinically evident basal cell carcinomas = the total number of target and/or non-target lesions present in individual subjects. Subjects in the Intent-to-Treat Analysis Population (defined as all randomized subjects) with available data were included in the analysis. The last observation carried forward method was used.	
End point type	Primary
End point timeframe: Baseline; Week 73	

End point values	Vismodegib Intermittent Schedule	Vismodegib Induction Followed by Intermittent Schedule		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	114	113		
Units: percent change				
arithmetic mean (standard deviation)	62.9 (± 52.01)	54.9 (± 54.85)		

### Statistical analyses

Statistical analysis title	Difference in mean relative reduction
Statistical analysis description: The mean difference in the mean relative reduction between treatment arms, along with the corresponding 95% confidence interval, was estimated by fitting an ANCOVA model with treatment as main effect and the following covariates: number of basal cell carcinomas at baseline, geographical region, immunosuppression status, confirmed basal cell carcinoma nevus syndrome. Asymptotic confidence intervals are presented for the difference between treatment arms.	
Comparison groups	Vismodegib Intermittent Schedule v Vismodegib Induction Followed by Intermittent Schedule

Number of subjects included in analysis	227
Analysis specification	Pre-specified
Analysis type	other
Parameter estimate	Mean difference (final values)
Point estimate	-8.3
Confidence interval	
level	95 %
sides	2-sided
lower limit	-22.2
upper limit	5.7

### Secondary: Percentage of Subjects Who Discontinued Study Treatment Due to Tolerability Issues

End point title	Percentage of Subjects Who Discontinued Study Treatment Due to Tolerability Issues
End point description:	The percentage of subjects who discontinued study treatment (due either to adverse event, refusal of treatment, or withdrawal of consent) was summarized by treatment group. Intent-to-Treat Analysis Population, defined as all randomized subjects.
End point type	Secondary
End point timeframe:	
Baseline to Week 73	

End point values	Vismodegib Intermittent Schedule	Vismodegib Induction Followed by Intermittent Schedule		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	116	113		
Units: percentage of subjects				
number (confidence interval 95%)				
Overall	37.1 (28.3 to 46.5)	41.6 (32.4 to 51.2)		
Adverse Events	20.7 (13.7 to 29.2)	27.4 (19.5 to 36.6)		
Refused Treatment	6 (2.5 to 12)	2.7 (0.6 to 7.6)		
Withdrew Consent	10.3 (5.5 to 17.4)	11.5 (6.3 to 18.9)		

### Statistical analyses

No statistical analyses for this end point

### Secondary: Mean Percent Change From Baseline in Total Size of Three Target Basal Cell Carcinoma Lesions in Individual Subjects at Week 73

End point title	Mean Percent Change From Baseline in Total Size of Three
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End point description:

The three target basal cell carcinoma lesions = the three largest visible lesions, at least 5 mm in the longest diameter, in individual subjects. Subjects in the Intent-to-Treat Analysis Population (defined as all randomized subjects) with available data were included in the analysis.

End point type Secondary

End point timeframe:

Baseline; Week 73

End point values	Vismodegib Intermittent Schedule	Vismodegib Induction Followed by Intermittent Schedule		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	94	85		
Units: percent change				
arithmetic mean (standard deviation)	82.9 (± 27.01)	68 (± 53.02)		

Statistical analyses

No statistical analyses for this end point

**Secondary: Percentage of Subjects With at Least 50% Reduction in the Number of Basal Cell Carcinomas at Week 73**

End point title Percentage of Subjects With at Least 50% Reduction in the Number of Basal Cell Carcinomas at Week 73

End point description:

Intent-to-Treat Analysis Population, defined as all randomized subjects.

End point type Secondary

End point timeframe:

Baseline; Week 73

End point values	Vismodegib Intermittent Schedule	Vismodegib Induction Followed by Intermittent Schedule		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	116	113		
Units: percentage of subjects				
number (not applicable)	65.5	50.4		

## Statistical analyses

No statistical analyses for this end point

### Secondary: Percentage of Subjects With New Basal Cell Carcinomas at Week 73

End point title	Percentage of Subjects With New Basal Cell Carcinomas at Week 73
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End point description:

Subjects in the Intent-to-Treat Analysis Population (defined as all randomized subjects) with available data were included in the analysis.

End point type	Secondary
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End point timeframe:

Baseline; Week 73

End point values	Vismodegib Intermittent Schedule	Vismodegib Induction Followed by Intermittent Schedule		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	94	86		
Units: percentage of subjects				
number (not applicable)				
No new lesions	76.6	74.4		
1 new lesion	10.6	11.6		
2 new lesions	5.3	5.8		
3 new lesions	5.3	2.3		
>3 new lesions	2.1	5.8		

## Statistical analyses

No statistical analyses for this end point

### Secondary: Percent Change in Total Number of Basal Cell Carcinomas Relative to Baseline at Week 85 (12 Weeks Following End of Treatment) (Recurrence Rate)

End point title	Percent Change in Total Number of Basal Cell Carcinomas Relative to Baseline at Week 85 (12 Weeks Following End of Treatment) (Recurrence Rate)
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End point description:

Subjects in the Intent-to-Treat Analysis Population (defined as all randomized subjects) with available data were included in the analysis.

End point type	Secondary
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End point timeframe:

Baseline; Week 85

End point values	Vismodegib Intermittent Schedule	Vismodegib Induction Followed by Intermittent Schedule		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	90	77		
Units: percent change				
arithmetic mean (standard deviation)	35.7 (± 50.25)	38.5 (± 55.22)		

### Statistical analyses

No statistical analyses for this end point

### Secondary: Percent Change in Total Number of Basal Cell Carcinomas Relative to Baseline at Week 97 (24 Weeks Following End of Treatment) (Recurrence Rate)

End point title	Percent Change in Total Number of Basal Cell Carcinomas Relative to Baseline at Week 97 (24 Weeks Following End of Treatment) (Recurrence Rate)
End point description: Subjects in the Intent-to-Treat Analysis Population (defined as all randomized subjects) with available data were included in the analysis.	
End point type	Secondary
End point timeframe: Baseline; Week 97	

End point values	Vismodegib Intermittent Schedule	Vismodegib Induction Followed by Intermittent Schedule		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	84	72		
Units: percent change				
arithmetic mean (standard deviation)	36 (± 49.48)	42.1 (± 57.83)		

### Statistical analyses

No statistical analyses for this end point

### Secondary: Percent Change in Total Number of Basal Cell Carcinomas Relative to Baseline at Week 125 (52 Weeks Following End of Treatment) (Recurrence Rate)

End point title	Percent Change in Total Number of Basal Cell Carcinomas Relative to Baseline at Week 125 (52 Weeks Following End of Treatment) (Recurrence Rate)
End point description: Subjects in the Intent-to-Treat Analysis Population (defined as all randomized subjects) with available data were included in the analysis.	

End point type	Secondary
End point timeframe:	
Baseline; Week 125	

End point values	Vismodegib Intermittent Schedule	Vismodegib Induction Followed by Intermittent Schedule		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	95	82		
Units: percent change				
arithmetic mean (standard deviation)	41.2 (± 45.23)	44 (± 42.87)		

### Statistical analyses

No statistical analyses for this end point

### Secondary: Percentage of Subjects Experiencing Any Adverse Event

End point title	Percentage of Subjects Experiencing Any Adverse Event
End point description:	
Safety Analysis Population: Subjects in the Intent-to-Treat Analysis Population (defined as all randomized subjects) who received at least one dose of study treatment.	
End point type	Secondary
End point timeframe:	
Up to 125 weeks	

End point values	Vismodegib Intermittent Schedule	Vismodegib Induction Followed by Intermittent Schedule		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	114	113		
Units: percentage of subjects				
number (not applicable)	99.1	97.3		

### Statistical analyses

No statistical analyses for this end point

### Secondary: Percent Change From Baseline in the Skindex-16 Symptom Domain Score at Week 73

End point title	Percent Change From Baseline in the Skindex-16 Symptom
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## End point description:

The Skindex-16 is a patient-reported outcome health questionnaire. Subjects were asked about their symptoms, and their answers were combined into a composite Symptom Domain Score. Scores range from 0 ("never bothered") to 100 ("always bothered"). Subjects in the Intent-to-Treat Analysis Population (defined as all randomized subjects) with available data were included in the analysis.

End point type	Secondary
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End point timeframe:

Baseline; Week 73

End point values	Vismodegib Intermittent Schedule	Vismodegib Induction Followed by Intermittent Schedule		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	87	79		
Units: percent change				
arithmetic mean (standard deviation)	-14.9 (± 25.75)	-12.6 (± 23.98)		

## Statistical analyses

No statistical analyses for this end point

**Secondary: Percent Change From Baseline in the Skindex-16 Emotion Domain Score at Week 73**

End point title	Percent Change From Baseline in the Skindex-16 Emotion Domain Score at Week 73
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## End point description:

The Skindex-16 is a patient-reported outcome health questionnaire. Subjects were asked about their emotional state, and their answers were combined into a composite Emotion Domain Score. Scores range from 0 ("never bothered") to 100 ("always bothered"). Subjects in the Intent-to-Treat Analysis Population (defined as all randomized subjects) with available data were included in the analysis.

End point type	Secondary
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End point timeframe:

Baseline; Week 73

End point values	Vismodegib Intermittent Schedule	Vismodegib Induction Followed by Intermittent Schedule		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	87	79		
Units: percent change				
arithmetic mean (standard deviation)	-27.4 (± 27.71)	-28.9 (± 28.16)		

## Statistical analyses

No statistical analyses for this end point

## Secondary: Percent Change From Baseline in the Skindex-16 Function Domain Score at Week 73

End point title	Percent Change From Baseline in the Skindex-16 Function Domain Score at Week 73
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End point description:

The Skindex-16 is a patient-reported outcome health questionnaire. Subjects were asked about their ability to function, and answers were combined into a composite Function Domain Score. Scores range from 0 ("never bothered") to 100 ("always bothered"). Subjects in the Intent-to-Treat Analysis Population (defined as all randomized subjects) with available data were included in the analysis.

End point type	Secondary
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End point timeframe:

Baseline; Week 73

End point values	Vismodegib Intermittent Schedule	Vismodegib Induction Followed by Intermittent Schedule		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	87	79		
Units: percent change				
arithmetic mean (standard deviation)	-9.5 (± 20.59)	-10.3 (± 26.03)		

## Statistical analyses

No statistical analyses for this end point

## Adverse events

### Adverse events information

Timeframe for reporting adverse events:

Up to 125 weeks

Adverse event reporting additional description:

Safety Analysis Population: Subjects in the Intent-to-Treat Analysis Population (defined as all randomized subjects) who received at least one dose of study treatment.

Assessment type	Systematic
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### Dictionary used

Dictionary name	MedDRA
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Dictionary version	18.0
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### Reporting groups

Reporting group title	Vismodegib Intermittent Schedule
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Reporting group description:

Vismodegib intermittent schedule of 12 weeks vismodegib followed by 8 weeks placebo, repeated 3 times with a final course of vismodegib (total 72 weeks), followed by 52 weeks treatment-free follow up

Reporting group title	Vismodegib Induction Followed by Intermittent Schedule
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Reporting group description:

Vismodegib beginning with 24 weeks induction followed by intermittent schedule 8 weeks placebo, 8 weeks vismodegib (total 72 weeks), followed by 52 weeks treatment-free follow up

Serious adverse events	Vismodegib Intermittent Schedule	Vismodegib Induction Followed by Intermittent Schedule	
Total subjects affected by serious adverse events			
subjects affected / exposed	24 / 114 (21.05%)	23 / 113 (20.35%)	
number of deaths (all causes)	2	2	
number of deaths resulting from adverse events			
Neoplasms benign, malignant and unspecified (incl cysts and polyps)			
Squamous cell carcinoma			
subjects affected / exposed	1 / 114 (0.88%)	2 / 113 (1.77%)	
occurrences causally related to treatment / all	0 / 1	0 / 2	
deaths causally related to treatment / all	0 / 0	0 / 0	
Metastatic squamous cell carcinoma			
subjects affected / exposed	1 / 114 (0.88%)	0 / 113 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Spindle cell sarcoma			

subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Squamous cell carcinoma of skin			
subjects affected / exposed	1 / 114 (0.88%)	0 / 113 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Seminoma			
subjects affected / exposed	1 / 114 (0.88%)	0 / 113 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Malignant melanoma in situ			
subjects affected / exposed	1 / 114 (0.88%)	0 / 113 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Vascular disorders			
Haematoma			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
General disorders and administration site conditions			
Asthenia			
subjects affected / exposed	1 / 114 (0.88%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	2 / 2	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Immune system disorders			
Primary amyloidosis			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Respiratory, thoracic and mediastinal disorders			
Pulmonary embolism			



subjects affected / exposed	1 / 114 (0.88%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	1 / 1	0 / 1	
deaths causally related to treatment / all	1 / 1	0 / 1	
Acute respiratory failure			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Chronic obstructive pulmonary disease			
subjects affected / exposed	1 / 114 (0.88%)	0 / 113 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Psychiatric disorders			
Depression			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Personality change			
subjects affected / exposed	1 / 114 (0.88%)	0 / 113 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Investigations			
Hepatic enzyme increased			
subjects affected / exposed	1 / 114 (0.88%)	0 / 113 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
International normalised ratio increased			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Platelet count decreased			
subjects affected / exposed	1 / 114 (0.88%)	0 / 113 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	

Injury, poisoning and procedural complications			
Ankle fracture			
subjects affected / exposed	1 / 114 (0.88%)	0 / 113 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hip fracture			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Humerus fracture			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Limb injury			
subjects affected / exposed	1 / 114 (0.88%)	0 / 113 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Thermal burn			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Arthropod bite			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Contusion			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Congenital, familial and genetic disorders			
Congenital cerebral cyst			

subjects affected / exposed	1 / 114 (0.88%)	0 / 113 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
<b>Cardiac disorders</b>			
Myocardial infarction			
subjects affected / exposed	1 / 114 (0.88%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 1	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Acute coronary syndrome			
subjects affected / exposed	1 / 114 (0.88%)	0 / 113 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Angina pectoris			
subjects affected / exposed	1 / 114 (0.88%)	0 / 113 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Bradycardia			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Cardiogenic shock			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 1	
Acute myocardial infarction			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
<b>Nervous system disorders</b>			
Cerebrovascular accident			
subjects affected / exposed	1 / 114 (0.88%)	0 / 113 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Lethargy			

subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	1 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Nervous system disorder			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Post herpetic neuralgia			
subjects affected / exposed	1 / 114 (0.88%)	0 / 113 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Tremor			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Transient ischaemic attack			
subjects affected / exposed	1 / 114 (0.88%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 1	0 / 2	
deaths causally related to treatment / all	0 / 0	0 / 0	
Blood and lymphatic system disorders			
Anaemia			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Febrile neutropenia			
subjects affected / exposed	1 / 114 (0.88%)	0 / 113 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pseudolymphoma			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	1 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Gastrointestinal disorders			

Abdominal pain			
subjects affected / exposed	1 / 114 (0.88%)	0 / 113 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Duodenal ulcer			
subjects affected / exposed	1 / 114 (0.88%)	0 / 113 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Gastric perforation			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pancreatitis acute			
subjects affected / exposed	1 / 114 (0.88%)	0 / 113 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hepatobiliary disorders			
Acute hepatic failure			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Skin and subcutaneous tissue disorders			
Xanthelasma			
subjects affected / exposed	1 / 114 (0.88%)	0 / 113 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Renal and urinary disorders			
Acute kidney injury			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 2	
deaths causally related to treatment / all	0 / 0	0 / 0	
Calculus urinary			

subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
<b>Musculoskeletal and connective tissue disorders</b>			
Arthralgia			
subjects affected / exposed	1 / 114 (0.88%)	0 / 113 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Back pain			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
<b>Infections and infestations</b>			
Pneumonia			
subjects affected / exposed	3 / 114 (2.63%)	0 / 113 (0.00%)	
occurrences causally related to treatment / all	0 / 3	0 / 0	
deaths causally related to treatment / all	0 / 1	0 / 0	
Abscess limb			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Cellulitis			
subjects affected / exposed	1 / 114 (0.88%)	0 / 113 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Diverticulitis			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Gastrointestinal infection			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	

Kidney infection			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pyelonephritis			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pyelonephritis acute			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Subcutaneous abscess			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Metabolism and nutrition disorders			
Dehydration			
subjects affected / exposed	0 / 114 (0.00%)	1 / 113 (0.88%)	
occurrences causally related to treatment / all	0 / 0	1 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	

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

<b>Non-serious adverse events</b>	Vismodegib Intermittent Schedule	Vismodegib Induction Followed by Intermittent Schedule	
Total subjects affected by non-serious adverse events			
subjects affected / exposed	107 / 114 (93.86%)	108 / 113 (95.58%)	
Investigations			
Weight decreased			
subjects affected / exposed	24 / 114 (21.05%)	21 / 113 (18.58%)	
occurrences (all)	31	23	
Blood creatine phosphokinase increased			

subjects affected / exposed occurrences (all)	11 / 114 (9.65%) 14	15 / 113 (13.27%) 24	
Alanine aminotransferase increased subjects affected / exposed occurrences (all)	7 / 114 (6.14%) 8	5 / 113 (4.42%) 5	
Vascular disorders Hypertension subjects affected / exposed occurrences (all)	2 / 114 (1.75%) 6	6 / 113 (5.31%) 11	
Nervous system disorders Dysgeusia subjects affected / exposed occurrences (all)	76 / 114 (66.67%) 123	75 / 113 (66.37%) 111	
Ageusia subjects affected / exposed occurrences (all)	13 / 114 (11.40%) 16	14 / 113 (12.39%) 18	
Headache subjects affected / exposed occurrences (all)	10 / 114 (8.77%) 10	14 / 113 (12.39%) 14	
General disorders and administration site conditions Fatigue subjects affected / exposed occurrences (all)	24 / 114 (21.05%) 30	26 / 113 (23.01%) 31	
Asthenia subjects affected / exposed occurrences (all)	15 / 114 (13.16%) 29	21 / 113 (18.58%) 31	
Pyrexia subjects affected / exposed occurrences (all)	1 / 114 (0.88%) 1	8 / 113 (7.08%) 9	
Gastrointestinal disorders Diarrhoea subjects affected / exposed occurrences (all)	20 / 114 (17.54%) 35	20 / 113 (17.70%) 24	
Nausea subjects affected / exposed occurrences (all)	23 / 114 (20.18%) 32	14 / 113 (12.39%) 25	



Abdominal pain subjects affected / exposed occurrences (all)	8 / 114 (7.02%) 10	12 / 113 (10.62%) 12	
Constipation subjects affected / exposed occurrences (all)	9 / 114 (7.89%) 11	8 / 113 (7.08%) 10	
Abdominal pain upper subjects affected / exposed occurrences (all)	9 / 114 (7.89%) 12	5 / 113 (4.42%) 6	
Vomiting subjects affected / exposed occurrences (all)	7 / 114 (6.14%) 8	4 / 113 (3.54%) 4	
Respiratory, thoracic and mediastinal disorders Cough subjects affected / exposed occurrences (all)	6 / 114 (5.26%) 7	5 / 113 (4.42%) 9	
Skin and subcutaneous tissue disorders Alopecia subjects affected / exposed occurrences (all)	72 / 114 (63.16%) 100	73 / 113 (64.60%) 93	
Pruritus subjects affected / exposed occurrences (all)	8 / 114 (7.02%) 13	13 / 113 (11.50%) 14	
Actinic keratosis subjects affected / exposed occurrences (all)	10 / 114 (8.77%) 13	8 / 113 (7.08%) 11	
Eczema subjects affected / exposed occurrences (all)	6 / 114 (5.26%) 7	0 / 113 (0.00%) 0	
Rash subjects affected / exposed occurrences (all)	3 / 114 (2.63%) 3	7 / 113 (6.19%) 7	
Seborrhoeic dermatitis subjects affected / exposed occurrences (all)	6 / 114 (5.26%) 8	2 / 113 (1.77%) 2	
Psychiatric disorders			

Insomina subjects affected / exposed occurrences (all)	5 / 114 (4.39%) 6	6 / 113 (5.31%) 6	
Musculoskeletal and connective tissue disorders Muscle spasms subjects affected / exposed occurrences (all)  Arthralgia subjects affected / exposed occurrences (all)  Myalgia subjects affected / exposed occurrences (all)  Back pain subjects affected / exposed occurrences (all)	83 / 114 (72.81%) 184  18 / 114 (15.79%) 25  18 / 114 (15.79%) 27  7 / 114 (6.14%) 8	94 / 113 (83.19%) 194  16 / 113 (14.16%) 25  12 / 113 (10.62%) 18  7 / 113 (6.19%) 8	
Infections and infestations Nasopharyngitis subjects affected / exposed occurrences (all)  Folliculitis subjects affected / exposed occurrences (all)  Bronchitis subjects affected / exposed occurrences (all)  Upper respiratory tract infection subjects affected / exposed occurrences (all)	7 / 114 (6.14%) 10  10 / 114 (8.77%) 13  5 / 114 (4.39%) 5  7 / 114 (6.14%) 13	12 / 113 (10.62%) 15  8 / 113 (7.08%) 10  8 / 113 (7.08%) 10  5 / 113 (4.42%) 5	
Metabolism and nutrition disorders Decreased appetite subjects affected / exposed occurrences (all)	21 / 114 (18.42%) 32	17 / 113 (15.04%) 20	

## More information

### Substantial protocol amendments (globally)

Were there any global substantial amendments to the protocol? Yes

Date	Amendment
27 May 2013	Changes included: additional guidance on amenorrhea/irregular menses; additional laboratory testing to include creatine kinase to further investigate muscle spasm events; additional guidance on Vismodegib in seminal fluid to ensure male subjects did not donate sperm during treatment and for 2 months after treatment; updated eligibility criteria to exclude 1) subjects known or suspected to abuse alcohol and 2) with known rare hereditary disturbance of galactose metabolism; amended drug interaction language to reflect updated core documents regarding possible interactions with metabolize ethinyl estradiol contraceptive steroids; alteration to statistical analysis measure of precision.
01 February 2016	In accordance with the revised Investigator's Brochure, the protocol and informed consent forms of all ongoing vismodegib studies recommending 7 months for pregnancy prevention (breast-feeding and blood donations) were amended to reflect new information on elimination of vismodegib from the body after treatment discontinuation. The population PK model for elimination was updated and finalized with additional PK data up to 12 months post-treatment obtained in Study MO25616 (STEVIE), as a result of this analysis the time period for females to avoid pregnancy, breast-feeding and for males and females blood donations was increased from 7 to 9 months after the last administration of vismodegib.

Notes:

### Interruptions (globally)

Were there any global interruptions to the trial? No

### Limitations and caveats

None reported