



Clinical trial results:

A randomized Phase II, 2-armed study in transplant ineligible (TI) patients with newly diagnosed multiple myeloma (NDMM) comparing Carfilzomib + Thalidomide + dexamethasone (KTd) versus Carfilzomib + Lenalidomide + dexamethasone (KRd) induction therapy with respect to response rates and investigating a Carfilzomib (K) monotherapy maintenance strategy

Summary

EudraCT number	2016-000475-24
Trial protocol	AT DE
Global end of trial date	28 March 2024

Results information

Result version number	v1 (current)
This version publication date	18 October 2024
First version publication date	18 October 2024

Trial information

Trial identification

Sponsor protocol code	AGMT_MM-2
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Additional study identifiers

ISRCTN number	-
ClinicalTrials.gov id (NCT number)	NCT02891811
WHO universal trial number (UTN)	-

Notes:

Sponsors

Sponsor organisation name	AGMT
Sponsor organisation address	Gentzgasse 60/21, Vienna, Austria, 1180
Public contact	Daniela Wolkersdorfer, AGMT, +43 6626404412, d.wolkersdofer@agmt.at
Scientific contact	Richard Greil, AGMT, +43 6626404412, office@agmt.at

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	24 March 2023
Is this the analysis of the primary completion data?	Yes
Primary completion date	24 March 2023
Global end of trial reached?	Yes
Global end of trial date	28 March 2024
Was the trial ended prematurely?	No

Notes:

General information about the trial

Main objective of the trial:

To show non-inferiority with respect to response rates between KTd and KRd: Overall response rate (ORR) are assessed according to International Myeloma Working Group (IMWG) criteria to determine the ORR in patients NDMM after receiving max 9 cycles induction therapy with either carfilzomib in combination with thalidomide and dexamethasone or carfilzomib in combination with lenalidomide and dexamethasone

Protection of trial subjects:

Anti-biotic, anti-viral, and anti-thrombosis pretreatment and prophylaxis was mandatory. Guidelines for dose modifications on case of toxicities were given. In general, concomitant medications and therapies necessary for supportive care and safety of the patient were allowed. The administration of any other anticancer agent or other concurrent investigational drug was not permitted. The inclusion of women of childbearing potential (WOCBP) and male subjects with pregnant or non-pregnant WOCBP had to follow specific recommendations for contraception and pregnancy testing. Furthermore, for all participating patients special counselling regarding use of thalidomide or lenalidomide had to be done prior to each treatment cycle.

Background therapy: -

Evidence for comparator: -

Actual start date of recruitment	20 March 2017
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	Austria: 121
Country: Number of subjects enrolled	Germany: 3
Worldwide total number of subjects	124
EEA total number of subjects	124

Notes:

Subjects enrolled per age group

In utero	0
Preterm newborn - gestational age < 37 wk	0
Newborns (0-27 days)	0
Infants and toddlers (28 days-23 months)	0

Children (2-11 years)	0
Adolescents (12-17 years)	0
Adults (18-64 years)	4
From 65 to 84 years	118
85 years and over	2

Subject disposition

Recruitment

Recruitment details:

Between 20-Mar-2017 and 16-Dec-2021, 124 patients were enrolled in 16 sites in Austria and 2 sites in Germany. 1 patient withdrew consent shortly after randomization, did not receive any study treatment and is excluded from analysis.

Pre-assignment

Screening details:

Adult patients (≥ 18 years) with newly diagnosed, symptomatic MM not eligible or not willing to undergo autologous stem cell transplantation following induction and who are $<$ NYHA class III/IV, ECOG PS 0-1, CrCl $>$ 30ml/min, PN ≤ 2 (without pain).

Period 1

Period 1 title	Induction
Is this the baseline period?	Yes
Allocation method	Randomised - controlled
Blinding used	Not blinded

Arms

Are arms mutually exclusive?	Yes
Arm title	KTd Induction

Arm description:

Induction therapy for a maximum of 9 cycles

Arm type	Experimental
Investigational medicinal product name	Carfilzomib
Investigational medicinal product code	
Other name	Kyprolis
Pharmaceutical forms	Powder for solution for infusion
Routes of administration	Infusion

Dosage and administration details:

Cycle 1: 20mg/m² on day 1+2, 27mg/m² on day 8, 9, 15 and 16; Cycle 2: 27mg/m² on day 1,2,8,9,15 and 16; Cycle 3-9: 56mg/m² on day 1, 8 and 15

Investigational medicinal product name	Thalidomide
Investigational medicinal product code	
Other name	
Pharmaceutical forms	Capsule, hard
Routes of administration	Oral use

Dosage and administration details:

100mg/day, day 1-28 (50mg in patients aged ≥ 75 years)

Investigational medicinal product name	Dexamethasone
Investigational medicinal product code	
Other name	
Pharmaceutical forms	Injection/infusion, Tablet
Routes of administration	Infusion , Injection , Oral use

Dosage and administration details:

40mg/week- day 1,8,15,22 (20mg in patients aged ≥ 75 years)

Arm title	KRd Induction
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Arm description:

Induction therapy for a maximum of 9 cycles

Arm type	Experimental
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Investigational medicinal product name	Carfilzomib
Investigational medicinal product code	
Other name	Kyprolis
Pharmaceutical forms	Powder for solution for infusion
Routes of administration	Infusion

Dosage and administration details:

Cycle 1: 20mg/m² on day 1+2, 27mg/m² on day 8, 9, 15 and 16; Cycle 2: 27mg/m² on day 1,2,8,9,15 and 16; Cycle 3-9: 56mg/m² on day 1, 8 and 15

Investigational medicinal product name	Lenalidomide
Investigational medicinal product code	
Other name	
Pharmaceutical forms	Capsule, hard
Routes of administration	Oral use

Dosage and administration details:

25mg/day, day 1-21

Investigational medicinal product name	Dexamethasone
Investigational medicinal product code	
Other name	
Pharmaceutical forms	Injection/infusion, Tablet
Routes of administration	Infusion , Injection , Oral use

Dosage and administration details:

40mg/week- day 1,8,15,22 (20mg in patients aged ≥75 years)

Number of subjects in period 1^[1]	KTd Induction	KRd Induction
Started	63	60
Completed	46	38
Not completed	17	22
investigator's decision	2	1
severe AE/toxicity	9	15
progressive disease/death	2	3
patient's decision	4	2
dialysis-dependent	-	1

Notes:

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

Justification: One patient withdrew consent shortly after randomization in KRd arm and did not receive any study treatment. This patient is excluded from analysis.

Period 2

Period 2 title	Maintenance
Is this the baseline period?	No
Allocation method	Randomised - controlled
Blinding used	Not blinded

Arms

Are arms mutually exclusive?	Yes
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Arm title	K monotherapy
Arm description: Maintenance treatment with carfilzomib for a maximum period of 12 months.	
Arm type	Experimental
Investigational medicinal product name	Carfilzomib
Investigational medicinal product code	
Other name	Kyprolis
Pharmaceutical forms	Powder for solution for infusion
Routes of administration	Infusion
Dosage and administration details: day 1 and day 15 using the last tolerated dose for 12 cycles	
Arm title	Observation
Arm description: Observation for 12 months	
Arm type	No intervention
No investigational medicinal product assigned in this arm	

Number of subjects in period 2	K monotherapy	Observation
Started	42	42
Completed	28	26
Not completed	14	16
investigator's decision	2	2
ongoing at primary endpoint analysis	1	1
progressive disease/death	7	13
severe AE/toxicity	2	-
patient's decision	2	-

Baseline characteristics

Reporting groups

Reporting group title	KTd Induction
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Reporting group description:

Induction therapy for a maximum of 9 cycles

Reporting group title	KRd Induction
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Reporting group description:

Induction therapy for a maximum of 9 cycles

Reporting group values	KTd Induction	KRd Induction	Total
Number of subjects	63	60	123
Age categorical Units: Subjects			
In utero			0
Preterm newborn infants (gestational age < 37 wks)			0
Newborns (0-27 days)			0
Infants and toddlers (28 days-23 months)			0
Children (2-11 years)			0
Adolescents (12-17 years)			0
Adults (18-64 years)			0
From 65-84 years			0
85 years and over			0
Age continuous Units: years			
median	75	75	
full range (min-max)	58 to 84	55 to 84	-
Gender categorical Units: Subjects			
Female	31	24	55
Male	32	36	68

End points

End points reporting groups

Reporting group title	KTd Induction
Reporting group description: Induction therapy for a maximum of 9 cycles	
Reporting group title	KRd Induction
Reporting group description: Induction therapy for a maximum of 9 cycles	
Reporting group title	K monotherapy
Reporting group description: Maintenance treatment with carfilzomib for a maximum period of 12 months.	
Reporting group title	Observation
Reporting group description: Observation for 12 months	

Primary: Response

End point title	Response
End point description: Response in patients treated with KTd or KRd during induction therapy. Patients who received study medication for at least 4 weeks were evaluable for response.	
End point type	Primary
End point timeframe: Response assessment was done at day 1 of each cycle (except cycle 1) and at the end of K combination treatment	

End point values	KTd Induction	KRd Induction		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	61	55		
Units: Subjects				
stringent complete response (sCR)	7	4		
complete response (CR)	20	17		
very good partial response (VGPR)	21	19		
partial response (PR)	8	8		
minimal response (MR)	3	2		
stable disease (SD)	2	4		
progressive disease (PD)	0	1		

Statistical analyses

Statistical analysis title	Overall response rates
Statistical analysis description: Response rates were similar during induction therapy with KTd (91.8%) and KRd (87.3%), meeting the requirements for non-inferiority.	

Comparison groups	KTd Induction v KRd Induction
Number of subjects included in analysis	116
Analysis specification	Pre-specified
Analysis type	non-inferiority ^[1]
Parameter estimate	Difference
Point estimate	-4.5
Confidence interval	
level	95 %
sides	2-sided
lower limit	-17
upper limit	7.1

Notes:

[1] - Non-inferiority is postulated if the CI of the differences does not cross 11%. CI was calculated using the Miettinen-Nurminen method

Adverse events

Adverse events information

Timeframe for reporting adverse events:

All patients having received at least one dose of the study medication were followed for adverse events for at least 28 days after discontinuing study treatment or completion of study treatment.

Assessment type	Systematic
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Dictionary used

Dictionary name	MedDRA
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Dictionary version	24.1
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Reporting groups

Reporting group title	Safety population
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Reporting group description:

The safety population includes all enrolled patients who received at least one dose of the study treatment. SARs are assessed as related to carfilzomib.

Serious adverse events	Safety population		
Total subjects affected by serious adverse events			
subjects affected / exposed	86 / 123 (69.92%)		
number of deaths (all causes)	23		
number of deaths resulting from adverse events	6		
Neoplasms benign, malignant and unspecified (incl cysts and polyps)			
Basal cell carcinoma			
subjects affected / exposed	2 / 123 (1.63%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
Myelodysplastic syndrome			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Bowen's disease			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Vascular disorders			
Hypotension			

subjects affected / exposed	2 / 123 (1.63%)			
occurrences causally related to treatment / all	1 / 2			
deaths causally related to treatment / all	0 / 0			
Embolism				
subjects affected / exposed	2 / 123 (1.63%)			
occurrences causally related to treatment / all	2 / 2			
deaths causally related to treatment / all	0 / 0			
Hypertensive crisis				
subjects affected / exposed	1 / 123 (0.81%)			
occurrences causally related to treatment / all	1 / 1			
deaths causally related to treatment / all	0 / 0			
Thrombophlebitis				
subjects affected / exposed	1 / 123 (0.81%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Circulatory collapse				
subjects affected / exposed	1 / 123 (0.81%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Thrombosis				
subjects affected / exposed	1 / 123 (0.81%)			
occurrences causally related to treatment / all	1 / 1			
deaths causally related to treatment / all	0 / 0			
Deep vein thrombosis				
subjects affected / exposed	1 / 123 (0.81%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Arterial stenosis				
subjects affected / exposed	1 / 123 (0.81%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Hypertension				

subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
General disorders and administration site conditions			
Pyrexia			
subjects affected / exposed	5 / 123 (4.07%)		
occurrences causally related to treatment / all	2 / 5		
deaths causally related to treatment / all	0 / 0		
Asthenia			
subjects affected / exposed	3 / 123 (2.44%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 0		
General physical health deterioration			
subjects affected / exposed	3 / 123 (2.44%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 0		
Pain			
subjects affected / exposed	2 / 123 (1.63%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
Death			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
Oedema			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Drug intolerance			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Infusion site reaction			

subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Hernia			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Reproductive system and breast disorders			
Cystocele			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Respiratory, thoracic and mediastinal disorders			
Pulmonary embolism			
subjects affected / exposed	2 / 123 (1.63%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 1		
Dyspnoea			
subjects affected / exposed	2 / 123 (1.63%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	0 / 0		
Pulmonary congestion			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Chronic obstructive pulmonary disease			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Pleural effusion			

subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
Psychiatric disorders			
Suicide attempt			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Confusional state			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Product issues			
Device dislocation			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Investigations			
C-reactive protein increased			
subjects affected / exposed	2 / 123 (1.63%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	0 / 0		
Hepatic enzyme increased			
subjects affected / exposed	2 / 123 (1.63%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	0 / 0		
Blood creatinine increased			
subjects affected / exposed	2 / 123 (1.63%)		
occurrences causally related to treatment / all	2 / 2		
deaths causally related to treatment / all	0 / 0		
Blood creatine increased			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		

Injury, poisoning and procedural complications			
Accidental overdose			
subjects affected / exposed	3 / 123 (2.44%)		
occurrences causally related to treatment / all	0 / 4		
deaths causally related to treatment / all	0 / 0		
Infusion related reaction			
subjects affected / exposed	2 / 123 (1.63%)		
occurrences causally related to treatment / all	2 / 2		
deaths causally related to treatment / all	0 / 0		
Thoracic vertebral fracture			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Fall			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Wound haemorrhage			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Ankle fracture			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Lumbar vertebral fracture			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Cardiac disorders			
Cardiac failure			
subjects affected / exposed	8 / 123 (6.50%)		
occurrences causally related to treatment / all	7 / 8		
deaths causally related to treatment / all	0 / 0		

Sinus bradycardia			
subjects affected / exposed	2 / 123 (1.63%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	0 / 0		
Atrial fibrillation			
subjects affected / exposed	2 / 123 (1.63%)		
occurrences causally related to treatment / all	2 / 2		
deaths causally related to treatment / all	0 / 0		
Tachycardia			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Angina pectoris			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Myocardial infarction			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Nervous system disorders			
Syncope			
subjects affected / exposed	2 / 123 (1.63%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 0		
Cerebrovascular accident			
subjects affected / exposed	2 / 123 (1.63%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 0		
Polyneuropathy			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Epilepsy			

subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Depressed level of consciousness			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Transient ischaemic attack			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Peroneal nerve palsy			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Facial paralysis			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Blood and lymphatic system disorders			
Anaemia			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Thrombotic microangiopathy			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	1 / 1		
Iron deficiency anaemia			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Gastrointestinal disorders			

Nausea				
subjects affected / exposed	3 / 123 (2.44%)			
occurrences causally related to treatment / all	2 / 4			
deaths causally related to treatment / all	0 / 0			
Vomiting				
subjects affected / exposed	3 / 123 (2.44%)			
occurrences causally related to treatment / all	2 / 3			
deaths causally related to treatment / all	0 / 0			
Enteritis				
subjects affected / exposed	2 / 123 (1.63%)			
occurrences causally related to treatment / all	1 / 2			
deaths causally related to treatment / all	0 / 0			
Haematochezia				
subjects affected / exposed	1 / 123 (0.81%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Rectal haemorrhage				
subjects affected / exposed	1 / 123 (0.81%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Gastritis				
subjects affected / exposed	1 / 123 (0.81%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Diarrhoea				
subjects affected / exposed	1 / 123 (0.81%)			
occurrences causally related to treatment / all	1 / 1			
deaths causally related to treatment / all	0 / 0			
Gastrointestinal haemorrhage				
subjects affected / exposed	1 / 123 (0.81%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Hepatobiliary disorders				

Hepatic failure			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Skin and subcutaneous tissue disorders			
Rash			
subjects affected / exposed	6 / 123 (4.88%)		
occurrences causally related to treatment / all	3 / 7		
deaths causally related to treatment / all	0 / 0		
Toxic epidermal necrolysis			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Rash maculo-papular			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Dermatitis acneiform			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Decubitus ulcer			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Drug reaction with eosinophilia and systemic symptoms			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Renal and urinary disorders			
Acute kidney injury			
subjects affected / exposed	4 / 123 (3.25%)		
occurrences causally related to treatment / all	0 / 4		
deaths causally related to treatment / all	0 / 1		

Urinary retention				
subjects affected / exposed	1 / 123 (0.81%)			
occurrences causally related to treatment / all	1 / 1			
deaths causally related to treatment / all	0 / 0			
Chronic kidney disease				
subjects affected / exposed	1 / 123 (0.81%)			
occurrences causally related to treatment / all	0 / 2			
deaths causally related to treatment / all	0 / 0			
Renal failure				
subjects affected / exposed	3 / 123 (2.44%)			
occurrences causally related to treatment / all	1 / 3			
deaths causally related to treatment / all	0 / 0			
Musculoskeletal and connective tissue disorders				
Spinal pain				
subjects affected / exposed	1 / 123 (0.81%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Neck pain				
subjects affected / exposed	1 / 123 (0.81%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Back pain				
subjects affected / exposed	1 / 123 (0.81%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Bone lesion				
subjects affected / exposed	1 / 123 (0.81%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Rheumatoid arthritis				
subjects affected / exposed	1 / 123 (0.81%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			

Bone pain			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Thoracic spinal stenosis			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Gouty arthritis			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Intervertebral disc disorder			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Infections and infestations			
Pneumonia			
subjects affected / exposed	15 / 123 (12.20%)		
occurrences causally related to treatment / all	11 / 16		
deaths causally related to treatment / all	0 / 0		
Sepsis			
subjects affected / exposed	4 / 123 (3.25%)		
occurrences causally related to treatment / all	2 / 4		
deaths causally related to treatment / all	0 / 0		
Urinary tract infection			
subjects affected / exposed	2 / 123 (1.63%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
Bronchitis			
subjects affected / exposed	2 / 123 (1.63%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
Upper respiratory tract infection			

subjects affected / exposed	2 / 123 (1.63%)			
occurrences causally related to treatment / all	1 / 2			
deaths causally related to treatment / all	0 / 0			
COVID-19				
subjects affected / exposed	2 / 123 (1.63%)			
occurrences causally related to treatment / all	0 / 2			
deaths causally related to treatment / all	0 / 0			
Diverticulitis				
subjects affected / exposed	2 / 123 (1.63%)			
occurrences causally related to treatment / all	1 / 2			
deaths causally related to treatment / all	0 / 0			
Influenza				
subjects affected / exposed	2 / 123 (1.63%)			
occurrences causally related to treatment / all	0 / 2			
deaths causally related to treatment / all	0 / 0			
Sinusitis				
subjects affected / exposed	1 / 123 (0.81%)			
occurrences causally related to treatment / all	1 / 1			
deaths causally related to treatment / all	0 / 0			
Respiratory tract infection				
subjects affected / exposed	1 / 123 (0.81%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Infection				
subjects affected / exposed	1 / 123 (0.81%)			
occurrences causally related to treatment / all	1 / 1			
deaths causally related to treatment / all	0 / 0			
Viral pericarditis				
subjects affected / exposed	1 / 123 (0.81%)			
occurrences causally related to treatment / all	1 / 1			
deaths causally related to treatment / all	1 / 1			
Febrile infection				

subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
Neuroborreliosis			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Gastroenteritis norovirus			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Osteomyelitis			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Viral myocarditis			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	1 / 1		
Escherichia bacteraemia			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Bacteraemia			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Mastoiditis			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Metabolism and nutrition disorders			
Hypocalcaemia			

subjects affected / exposed	2 / 123 (1.63%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 0		
Cachexia			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
Hypokalaemia			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Decreased appetite			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Hypercalcaemia			
subjects affected / exposed	1 / 123 (0.81%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		

Frequency threshold for reporting non-serious adverse events: 4 %

Non-serious adverse events	Safety population		
Total subjects affected by non-serious adverse events			
subjects affected / exposed	118 / 123 (95.93%)		
Vascular disorders			
Hypertension			
subjects affected / exposed	12 / 123 (9.76%)		
occurrences (all)	20		
Hypotension			
subjects affected / exposed	8 / 123 (6.50%)		
occurrences (all)	8		
Thrombophlebitis			

subjects affected / exposed	6 / 123 (4.88%)		
occurrences (all)	7		
General disorders and administration site conditions			
Fatigue			
subjects affected / exposed	63 / 123 (51.22%)		
occurrences (all)	110		
Oedema peripheral			
subjects affected / exposed	44 / 123 (35.77%)		
occurrences (all)	71		
Pyrexia			
subjects affected / exposed	19 / 123 (15.45%)		
occurrences (all)	29		
Pain			
subjects affected / exposed	12 / 123 (9.76%)		
occurrences (all)	18		
Chills			
subjects affected / exposed	8 / 123 (6.50%)		
occurrences (all)	9		
Asthenia			
subjects affected / exposed	8 / 123 (6.50%)		
occurrences (all)	9		
Chest pain			
subjects affected / exposed	7 / 123 (5.69%)		
occurrences (all)	7		
Influenza like illness			
subjects affected / exposed	7 / 123 (5.69%)		
occurrences (all)	7		
Oedema			
subjects affected / exposed	6 / 123 (4.88%)		
occurrences (all)	7		
Respiratory, thoracic and mediastinal disorders			
Dyspnoea			
subjects affected / exposed	23 / 123 (18.70%)		
occurrences (all)	29		
Cough			

subjects affected / exposed occurrences (all) Oropharyngeal pain subjects affected / exposed occurrences (all)	13 / 123 (10.57%) 15 10 / 123 (8.13%) 10		
Psychiatric disorders Sleep disorder subjects affected / exposed occurrences (all) Insomnia subjects affected / exposed occurrences (all)	10 / 123 (8.13%) 11 9 / 123 (7.32%) 12		
Investigations C-reactive protein increased subjects affected / exposed occurrences (all) Platelet count decreased subjects affected / exposed occurrences (all) Neutrophil count decreased subjects affected / exposed occurrences (all)	19 / 123 (15.45%) 22 8 / 123 (6.50%) 27 7 / 123 (5.69%) 18		
Nervous system disorders Polyneuropathy subjects affected / exposed occurrences (all) Tremor subjects affected / exposed occurrences (all) Paraesthesia subjects affected / exposed occurrences (all) Neuropathy peripheral subjects affected / exposed occurrences (all) Headache	22 / 123 (17.89%) 27 15 / 123 (12.20%) 17 11 / 123 (8.94%) 12 8 / 123 (6.50%) 9		

subjects affected / exposed	14 / 123 (11.38%)		
occurrences (all)	16		
Blood and lymphatic system disorders			
Anaemia			
subjects affected / exposed	20 / 123 (16.26%)		
occurrences (all)	33		
Thrombocytopenia			
subjects affected / exposed	17 / 123 (13.82%)		
occurrences (all)	34		
Neutropenia			
subjects affected / exposed	11 / 123 (8.94%)		
occurrences (all)	17		
Leukopenia			
subjects affected / exposed	6 / 123 (4.88%)		
occurrences (all)	8		
Ear and labyrinth disorders			
Vertigo			
subjects affected / exposed	27 / 123 (21.95%)		
occurrences (all)	35		
Tinnitus			
subjects affected / exposed	7 / 123 (5.69%)		
occurrences (all)	9		
Eye disorders			
Visual impairment			
subjects affected / exposed	6 / 123 (4.88%)		
occurrences (all)	7		
Cataract			
subjects affected / exposed	6 / 123 (4.88%)		
occurrences (all)	6		
Gastrointestinal disorders			
Nausea			
subjects affected / exposed	37 / 123 (30.08%)		
occurrences (all)	67		
Constipation			
subjects affected / exposed	32 / 123 (26.02%)		
occurrences (all)	40		
Diarrhoea			

subjects affected / exposed	27 / 123 (21.95%)		
occurrences (all)	43		
Vomiting			
subjects affected / exposed	14 / 123 (11.38%)		
occurrences (all)	21		
Abdominal pain upper			
subjects affected / exposed	6 / 123 (4.88%)		
occurrences (all)	7		
Dry mouth			
subjects affected / exposed	6 / 123 (4.88%)		
occurrences (all)	7		
Skin and subcutaneous tissue disorders			
Rash			
subjects affected / exposed	25 / 123 (20.33%)		
occurrences (all)	39		
Pruritus			
subjects affected / exposed	9 / 123 (7.32%)		
occurrences (all)	11		
Musculoskeletal and connective tissue disorders			
Back pain			
subjects affected / exposed	18 / 123 (14.63%)		
occurrences (all)	22		
Arthralgia			
subjects affected / exposed	18 / 123 (14.63%)		
occurrences (all)	25		
Muscle spasms			
subjects affected / exposed	17 / 123 (13.82%)		
occurrences (all)	27		
Bone pain			
subjects affected / exposed	10 / 123 (8.13%)		
occurrences (all)	15		
Pain in extremity			
subjects affected / exposed	9 / 123 (7.32%)		
occurrences (all)	12		
Myalgia			

subjects affected / exposed occurrences (all)	7 / 123 (5.69%) 10		
Infections and infestations			
Nasopharyngitis			
subjects affected / exposed	19 / 123 (15.45%)		
occurrences (all)	20		
Urinary tract infection			
subjects affected / exposed	14 / 123 (11.38%)		
occurrences (all)	15		
Bronchitis			
subjects affected / exposed	11 / 123 (8.94%)		
occurrences (all)	14		
Infection			
subjects affected / exposed	8 / 123 (6.50%)		
occurrences (all)	8		
Metabolism and nutrition disorders			
Decreased appetite			
subjects affected / exposed	17 / 123 (13.82%)		
occurrences (all)	19		
Hypokalaemia			
subjects affected / exposed	9 / 123 (7.32%)		
occurrences (all)	9		
Hypocalcaemia			
subjects affected / exposed	7 / 123 (5.69%)		
occurrences (all)	11		

More information

Substantial protocol amendments (globally)

Were there any global substantial amendments to the protocol? Yes

Date	Amendment
31 January 2017	Based on data from Biran et al. 2016, presented at ASH 2016, increased cardiotoxicity and thrombotic microangiopathy were seen in elderly patients receiving 70 mg carfilzomib in combination with dexamethasone and lenalidomide. Due to these observations, the dose and scheduling of carfilzomib was adapted for this trial. Also frail patients and those with an ECOG PS status ≥ 2 were excluded from this study.

Notes:

Interruptions (globally)

Were there any global interruptions to the trial? No

Limitations and caveats

None reported

Online references

<http://www.ncbi.nlm.nih.gov/pubmed/38895059>

<http://www.ncbi.nlm.nih.gov/pubmed/38425185>