



## Clinical trial results:

**A multicenter, phase II study of bortezomib and dexamethasone as induction treatment followed by high dose melphalan (HDM) and autologous stem cell transplantation (SCT) in patients with de novo amyloid light chain (AL) amyloidosis.**

### Summary

EudraCT number	2010-021445-42
Trial protocol	NL BE DE
Global end of trial date	29 January 2021

### Results information

Result version number	v1 (current)
This version publication date	04 September 2022
First version publication date	04 September 2022

### Trial information

#### Trial identification

Sponsor protocol code	HO104
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#### Additional study identifiers

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

Notes:

### Sponsors

Sponsor organisation name	HOVON
Sponsor organisation address	De Boelelaan 1117, Amsterdam, Netherlands,
Public contact	HOVON Data Center, HOVON, 31 (0)107041560, hdc@erasmusmc.nl
Scientific contact	HOVON Data Center, HOVON, 31 (0)107041560, hdc@erasmusmc.nl

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	14 July 2017
Is this the analysis of the primary completion data?	Yes
Primary completion date	16 June 2016
Global end of trial reached?	Yes
Global end of trial date	29 January 2021
Was the trial ended prematurely?	No

Notes:

## General information about the trial

Main objective of the trial:

To determine the efficacy of bortezomib plus dexamethasone followed by HDM and auto-SCT in patients with newly diagnosed AL amyloidosis who are 18-70 years inclusive.

Protection of trial subjects:

Monitoring and Insurance

Background therapy: -

Evidence for comparator: -

Actual start date of recruitment	12 January 2012
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: 45
Country: Number of subjects enrolled	Belgium: 4
Country: Number of subjects enrolled	Germany: 8
Worldwide total number of subjects	57
EEA total number of subjects	57

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)	50
From 65 to 84 years	7
85 years and over	0

## Subject disposition

### Recruitment

Recruitment details: -

### Pre-assignment

Screening details:

All subjects gave written informed consent and were screened according to the inclusion- and exclusion criteria

### Period 1

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

### Arms

<b>Arm title</b>	Experimental group
Arm description: -	
Arm type	Experimental
Investigational medicinal product name	Bortezomib
Investigational medicinal product code	
Other name	
Pharmaceutical forms	Solution for injection
Routes of administration	Subcutaneous use

Dosage and administration details:

Patients will be treated with 4 cycles of bortezomib and dexamethasone (q 3 weeks). The recommended starting dose of bortezomib is 1.3 mg/m<sup>2</sup> and it is given subcutaneously in the hospital on an outpatient basis.

<b>Number of subjects in period 1</b>	Experimental group
Started	57
Completed	38
Not completed	19
Consent withdrawn by subject	1
not eligible for ASCT or other causes	9
Adverse events	5
Lack of efficacy	4

## Baseline characteristics

### Reporting groups

Reporting group title	Overall period
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Reporting group description: -

Reporting group values	Overall period	Total	
Number of subjects	57	57	
Age categorical			
Units: Subjects			
In utero	0	0	
Preterm newborn infants (gestational age < 37 wks)	0	0	
Newborns (0-27 days)	0	0	
Infants and toddlers (28 days-23 months)	0	0	
Children (2-11 years)	0	0	
Adolescents (12-17 years)	0	0	
Adults (18-64 years)	0	0	
From 65-84 years	0	0	
85 years and over	0	0	
Adults (18-70 years)	57	57	
Age continuous			
Units: years			
median	59		
full range (min-max)	26 to 70	-	
Gender categorical			
Units: Subjects			
Female	25	25	
Male	32	32	

## End points

### End points reporting groups

Reporting group title	Experimental group
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Reporting group description: -

### Primary: Primary endpoint

End point title	Primary endpoint <sup>[1]</sup>
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End point description:

End point type	Primary
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End point timeframe:

See publication.

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: see attached chart/documents for results

<b>End point values</b>	Experimental group			
Subject group type	Reporting group			
Number of subjects analysed	57			
Units: Whole	57			

### Attachments (see zip file)

Statistical data section from publication/HO104\_statistical  
List of reported non-SAE's/nonsaedata104-7Jul2022.pdf  
List of reported SAE's/saedata104-7Jul2022.pdf

### Statistical analyses

No statistical analyses for this end point

## Adverse events

### Adverse events information

Timeframe for reporting adverse events:

Adverse events will be reported from the first study-related procedure until 30 days following the last dose of any drug from the protocol treatment schedule or until the start of subsequent systemic therapy for the disease under study, if earlier.

Adverse event reporting additional description:

Adverse events occurring after 30 days should also be reported if considered at least possibly related to the investigational medicinal product by the investigator.

Assessment type	Systematic
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### Dictionary used

Dictionary name	CTCAE
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Dictionary version	4
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### Reporting groups

Reporting group title	Experimental group
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Reporting group description: -

<b>Serious adverse events</b>	Experimental group		
Total subjects affected by serious adverse events			
subjects affected / exposed	35 / 57 (61.40%)		
number of deaths (all causes)	14		
number of deaths resulting from adverse events			
Vascular disorders			
Hypotension			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Orthostatic hypotension			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Symptomatic hypotension			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
General disorders and administration site conditions			
Oedema			

subjects affected / exposed	2 / 57 (3.51%)		
occurrences causally related to treatment / all	2 / 2		
deaths causally related to treatment / all	0 / 0		
General malaise			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Respiratory, thoracic and mediastinal disorders			
Dyspnea/ bronchial hyperreactivity			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Pulmonary bleeding			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Pulmonary embolism			
subjects affected / exposed	2 / 57 (3.51%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	0 / 0		
Injury, poisoning and procedural complications			
Sinusitis			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Cardiac disorders			
Cardiac arrest			
subjects affected / exposed	2 / 57 (3.51%)		
occurrences causally related to treatment / all	2 / 2		
deaths causally related to treatment / all	2 / 2		
Heart failure			

subjects affected / exposed	5 / 57 (8.77%)		
occurrences causally related to treatment / all	4 / 6		
deaths causally related to treatment / all	0 / 0		
Cardiac right ventricular dysfunction			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Complete atrioventricular block			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Myocardial infarction			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Nervous system disorders			
Intercranial hemorrhage			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Near syncope			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Poly neuropathy			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Blood and lymphatic system disorders			
Thrombopenia			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Gastrointestinal disorders			

Colitis			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Diarrhea			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Gastro–intestinal necrosis			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Ischemic colitis			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Pancreatitis			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Hepatobiliary disorders			
Cholecystitis			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Hepato–Renal syndrome			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
Renal and urinary disorders			
Acute kidney injury			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		

Acute tubulus necrosis			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Progress. of renal insufficiency			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Renal insufficiency			
subjects affected / exposed	2 / 57 (3.51%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
Musculoskeletal and connective tissue disorders			
Muscle weakness arms and legs			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Infections and infestations			
Community aquired pneumonia			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Gastrointestinal infection			
subjects affected / exposed	2 / 57 (3.51%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
Lung infection			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Pneumonia			
subjects affected / exposed	4 / 57 (7.02%)		
occurrences causally related to treatment / all	3 / 4		
deaths causally related to treatment / all	0 / 0		

Pneumosepsis			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Sepsis			
subjects affected / exposed	3 / 57 (5.26%)		
occurrences causally related to treatment / all	3 / 3		
deaths causally related to treatment / all	0 / 0		
Viral infection			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Metabolism and nutrition disorders			
Dehydration			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	2 / 2		
deaths causally related to treatment / all	0 / 0		
Fluid overflow			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Fluid retention			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Hypoglycemia			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Hyponatremia			
subjects affected / exposed	1 / 57 (1.75%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		

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

<b>Non-serious adverse events</b>	Experimental group		
Total subjects affected by non-serious adverse events subjects affected / exposed	52 / 57 (91.23%)		
Neoplasms benign, malignant and unspecified (incl cysts and polyps) Neoplasms benign, malignant and unspecified subjects affected / exposed occurrences (all)	2 / 57 (3.51%) 2	Additional description: All combined, see non-SAE chart for details	
Vascular disorders Vascular disorders subjects affected / exposed occurrences (all)	15 / 57 (26.32%) 20	Additional description: All combined, see non-SAE chart for details	
General disorders and administration site conditions General disorders and administration site conditions subjects affected / exposed occurrences (all)	32 / 57 (56.14%) 53	Additional description: All combined, see non-SAE chart for details	
Immune system disorders Immune system disorders subjects affected / exposed occurrences (all)	1 / 57 (1.75%) 1	Additional description: All combined, see non-SAE chart for details	
Reproductive system and breast disorders Reproductive system and breast disorders subjects affected / exposed occurrences (all)	1 / 57 (1.75%) 1	Additional description: All combined, see non-SAE chart for details	
Respiratory, thoracic and mediastinal disorders Respiratory, thoracic and mediastinal disorders subjects affected / exposed occurrences (all)	17 / 57 (29.82%) 20	Additional description: All combined, see non-SAE chart for details	
Psychiatric disorders Psychiatric disorders subjects affected / exposed occurrences (all)	12 / 57 (21.05%) 14	Additional description: All combined, see non-SAE chart for details	
Investigations			

Investigations subjects affected / exposed occurrences (all)	Additional description: All combined, see non-SAE chart for details		
	25 / 57 (43.86%)		
	50		
Injury, poisoning and procedural complications Injury, poisoning and procedural complications	Additional description: All combined, see non-SAE chart for details		
subjects affected / exposed	1 / 57 (1.75%)		
occurrences (all)	1		
Cardiac disorders Cardiac disorders	Additional description: All combined, see non-SAE chart for details		
subjects affected / exposed	13 / 57 (22.81%)		
occurrences (all)	21		
Nervous system disorders Nervous system disorders	Additional description: All combined, see non-SAE chart for details		
subjects affected / exposed	33 / 57 (57.89%)		
occurrences (all)	63		
Blood and lymphatic system disorders Blood and lymphatic disorders	Additional description: All combined, see non-SAE chart for details		
subjects affected / exposed	16 / 57 (28.07%)		
occurrences (all)	24		
Eye disorders Eye disorders	Additional description: All combined, see non-SAE chart for details		
subjects affected / exposed	5 / 57 (8.77%)		
occurrences (all)	8		
Gastrointestinal disorders Gastrointestinal disorders	Additional description: All combined, see non-SAE chart for details		
subjects affected / exposed	33 / 57 (57.89%)		
occurrences (all)	78		
Hepatobiliary disorders Hepatobiliary disorders	Additional description: All combined, see non-SAE chart for details		
subjects affected / exposed	2 / 57 (3.51%)		
occurrences (all)	2		
Skin and subcutaneous tissue disorders Skin and subcutaneous tissue disorders	Additional description: All combined, see non-SAE chart for details		
subjects affected / exposed	12 / 57 (21.05%)		
occurrences (all)	19		
Renal and urinary disorders			

Renal and urinary disorders subjects affected / exposed occurrences (all)	Additional description: All combined, see non-SAE chart for details		
	6 / 57 (10.53%) 8		
Musculoskeletal and connective tissue disorders Musculoskeletal and connective tissue disorders subjects affected / exposed occurrences (all)	Additional description: All combined, see non-SAE chart for details		
	11 / 57 (19.30%) 11		
Infections and infestations Infections and infestations subjects affected / exposed occurrences (all)	Additional description: All combined, see non-SAE chart for details		
	25 / 57 (43.86%) 40		
Metabolism and nutrition disorders Metabolism and nutrition disorders subjects affected / exposed occurrences (all)	Additional description: All combined, see non-SAE chart for details		
	27 / 57 (47.37%) 63		

## More information

### Substantial protocol amendments (globally)

Were there any global substantial amendments to the protocol? Yes

Date	Amendment
20 December 2011	AM1: change of administration route bortezomib (s.c. instead of i.v.). New version of protocol, ICF, bortezomib IB and bortezomib labels.
10 July 2012	AM2: addition of pre-study ICF, new version of protocol, ICF and bortezomib IB.
10 January 2013	AM3: new version of protocol, ICF and bortezomib IB.
27 March 2014	AM4: closure of dexamethasone-only arm, new version of protocol, ICF and bortezomib IB
17 July 2015	AM5: new version of protocol, ICF and bortezomib IB.

Notes:

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### Interruptions (globally)

Were there any global interruptions to the trial? No

### Limitations and caveats

None reported

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### Online references

<http://www.ncbi.nlm.nih.gov/pubmed/30923094>