



## Clinical trial results:

### A Randomised, Double-Blind, Placebo-Controlled, Dose-Response Study of the Efficacy and Safety of MEDI7352 in Subjects with Painful Diabetic Neuropathy

#### Summary

EudraCT number	2018-002523-42
Trial protocol	GB ES PL HU DK RO
Global end of trial date	29 June 2023

#### Results information

Result version number	v2 (current)
This version publication date	31 August 2024
First version publication date	14 July 2024
Version creation reason	

#### Trial information

##### Trial identification

Sponsor protocol code	D5680C00002
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##### Additional study identifiers

ISRCTN number	-
ClinicalTrials.gov id (NCT number)	NCT03755934
WHO universal trial number (UTN)	-

Notes:

#### Sponsors

Sponsor organisation name	AstraZeneca
Sponsor organisation address	151 85, Sodertalje, Sweden,
Public contact	Global Clinical Lead, AstraZeneca Clinical Study Information Center, +1 8772409479, information.center@astrazeneca.com
Scientific contact	Global Clinical Lead, AstraZeneca Clinical Study Information Center, +1 8772409479, information.center@astrazeneca.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 2023
Is this the analysis of the primary completion data?	No

Global end of trial reached?	Yes
Global end of trial date	29 June 2023
Was the trial ended prematurely?	Yes

Notes:

## General information about the trial

Main objective of the trial:

The primary objective of the study is to assess the efficacy of MEDI7352 versus placebo on chronic pain in subjects with painful diabetic neuropathy (PDN) currently taking standard of care medication for their PDN pain.

Protection of trial subjects:

The conduct of this clinical study met all local and regulatory requirements. The study was conducted in accordance with the ethical principles that have their origin in the Declaration of Helsinki and are consistent with International Conference on Harmonization guideline: Good Clinical Practice, and applicable regulatory requirements. Participants signed an informed consent form and could withdraw from the study at any time without any disadvantage and without having to provide a reason for this decision. Only investigators qualified by training and experience were selected as appropriate experts to investigate the study drug.

Background therapy: -

Evidence for comparator: -

Actual start date of recruitment	19 November 2018
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	Poland: 62
Country: Number of subjects enrolled	Romania: 6
Country: Number of subjects enrolled	United Kingdom: 34
Country: Number of subjects enrolled	Hungary: 10
Worldwide total number of subjects	112
EEA total number of subjects	78

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)	68
From 65 to 84 years	44
85 years and over	0

## Subject disposition

### Recruitment

Recruitment details:

The study was conducted at 45 sites in 4 countries (the United Kingdom, Hungary, Poland, and Romania).

### Pre-assignment

Screening details:

A total of 112 participants were randomized, of which 107 participants received at least one dose of study drug.

### 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, Carer

### Arms

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

Arm description:

Participants received intravenous (IV) placebo infusion matched to MEDI7352 during 12-week treatment period.

Arm type	Placebo
Investigational medicinal product name	Placebo
Investigational medicinal product code	
Other name	
Pharmaceutical forms	Solution for infusion
Routes of administration	Intravenous use

Dosage and administration details:

IV placebo infusion matched to MEDI7352 during 12-week treatment period.

<b>Arm title</b>	MEDI7352 Low Dose
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Arm description:

Participants received IV MEDI7352 infusion during 12-week treatment period.

Arm type	Experimental
Investigational medicinal product name	MEDI7352
Investigational medicinal product code	
Other name	
Pharmaceutical forms	Solution for infusion
Routes of administration	Intravenous use

Dosage and administration details:

IV MEDI7352 given during a 12-week treatment period.

<b>Arm title</b>	MEDI7352 Medium Dose
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Arm description:

Participants received IV MEDI7352 infusion during 12-week treatment period.

Arm type	Experimental
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Investigational medicinal product name	MEDI7352
Investigational medicinal product code	
Other name	
Pharmaceutical forms	Solution for infusion
Routes of administration	Intravenous use
Dosage and administration details: IV MEDI7352 given during a 12-week treatment period.	
<b>Arm title</b>	MEDI7352 High Dose

Arm description:

Participants received IV MEDI7352 infusion during 12-week treatment period.

Arm type	Experimental
Investigational medicinal product name	MEDI7352
Investigational medicinal product code	
Other name	
Pharmaceutical forms	Solution for infusion
Routes of administration	Intravenous use

Dosage and administration details:

IV MEDI7352 given during a 12-week treatment period.

<b>Number of subjects in period 1</b>	Placebo	MEDI7352 Low Dose	MEDI7352 Medium Dose
Started	54	6	16
Treated	54	4	14
Completed	40	4	12
Not completed	14	2	4
Physician decision	2	-	-
Consent withdrawn by subject	5	-	-
Adverse event, non-fatal	4	1	1
Unspecified	3	1	3

<b>Number of subjects in period 1</b>	MEDI7352 High Dose
Started	36
Treated	35
Completed	30
Not completed	6
Physician decision	1
Consent withdrawn by subject	1
Adverse event, non-fatal	3
Unspecified	1

## Baseline characteristics

### Reporting groups

Reporting group title	Placebo
Reporting group description:	
Participants received intravenous (IV) placebo infusion matched to MEDI7352 during 12-week treatment period.	
Reporting group title	MEDI7352 Low Dose
Reporting group description:	
Participants received IV MEDI7352 infusion during 12-week treatment period.	
Reporting group title	MEDI7352 Medium Dose
Reporting group description:	
Participants received IV MEDI7352 infusion during 12-week treatment period.	
Reporting group title	MEDI7352 High Dose
Reporting group description:	
Participants received IV MEDI7352 infusion during 12-week treatment period.	

Reporting group values	Placebo	MEDI7352 Low Dose	MEDI7352 Medium Dose
Number of subjects	54	6	16
Age Categorical Units: Participants			
In utero	0	0	0
Preterm newborn infants (gestational age < 37 wks)	0	0	0
Newborns (0-27 days)	0	0	0
Infants and toddlers (28 days-23 months)	0	0	0
Children (2-11 years)	0	0	0
Adolescents (12-17 years)	0	0	0
Adults (18-64 years)	35	3	9
From 65-84 years	19	3	7
85 years and over	0	0	0
Age continuous Units: years			
arithmetic mean	60.7	60.7	60.1
standard deviation	± 8.02	± 17.14	± 11.49
Sex: Female, Male Units: Participants			
Female	20	0	5
Male	34	6	11
Race (NIH/OMB) Units: Subjects			
American Indian or Alaska Native	0	0	0
Asian	0	0	0
Native Hawaiian or Other Pacific Islander	0	0	0
Black or African American	0	0	2
White	53	6	14
More than one race	0	0	0
Unknown or Not Reported	1	0	0

Ethnicity (NIH/OMB)			
Units: Subjects			
Hispanic or Latino	0	0	0
Not Hispanic or Latino	54	6	16
Unknown or Not Reported	0	0	0

Reporting group values	MEDI7352 High Dose	Total	
Number of subjects	36	112	
Age Categorical			
Units: Participants			
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)	21	68	
From 65-84 years	15	44	
85 years and over	0	0	
Age continuous			
Units: years			
arithmetic mean	60.1		
standard deviation	± 10.21	-	
Sex: Female, Male			
Units: Participants			
Female	15	40	
Male	21	72	
Race (NIH/OMB)			
Units: Subjects			
American Indian or Alaska Native	0	0	
Asian	0	0	
Native Hawaiian or Other Pacific Islander	0	0	
Black or African American	0	2	
White	36	109	
More than one race	0	0	
Unknown or Not Reported	0	1	
Ethnicity (NIH/OMB)			
Units: Subjects			
Hispanic or Latino	0	0	
Not Hispanic or Latino	36	112	
Unknown or Not Reported	0	0	

## End points

### End points reporting groups

Reporting group title	Placebo
Reporting group description: Participants received intravenous (IV) placebo infusion matched to MEDI7352 during 12-week treatment period.	
Reporting group title	MEDI7352 Low Dose
Reporting group description: Participants received IV MEDI7352 infusion during 12-week treatment period.	
Reporting group title	MEDI7352 Medium Dose
Reporting group description: Participants received IV MEDI7352 infusion during 12-week treatment period.	
Reporting group title	MEDI7352 High Dose
Reporting group description: Participants received IV MEDI7352 infusion during 12-week treatment period.	

### Primary: Change From Baseline in Weekly Average of Average Daily Pain Score to Week 12

End point title	Change From Baseline in Weekly Average of Average Daily Pain Score to Week 12
End point description: Change from baseline to Week 12 in weekly average of average daily pain score is reported. Participants assessed their perceived daily average neuropathic pain over the previous 24 hours using an 11-point numerical rating scale (NRS), with 0 representing no pain and 10 representing the worst pain imaginable. Participants were instructed to assess their average daily pain at approximately the same time every morning, and to record the response in a subject diary (electronic patient-reported outcome [ePRO]). Modified intent-to-treat (mITT) population included all participants who received at least 1 dose of any double-blind study drug and have at least 1 daily NRS assessment while receiving the treatment.	
End point type	Primary
End point timeframe: Baseline (Day -7 to Day -1, inclusive) through Week 12	

End point values	Placebo	MEDI7352 Low Dose	MEDI7352 Medium Dose	MEDI7352 High Dose
Subject group type	Reporting group	Reporting group	Reporting group	Reporting group
Number of subjects analysed	54	4	14	35
Units: Units on a scale				
arithmetic mean (standard deviation)	-1.244 (± 1.7327)	-3.387 (± 1.9605)	-0.615 (± 1.0431)	-2.699 (± 2.0819)

### Statistical analyses

Statistical analysis title	ANCOVA Analysis
Comparison groups	Placebo v MEDI7352 Low Dose



Number of subjects included in analysis	58
Analysis specification	Pre-specified
Analysis type	other
P-value	= 0.0437 <sup>[1]</sup>
Method	ANCOVA
Parameter estimate	LS mean estimate
Point estimate	-1.96
Confidence interval	
level	95 %
sides	2-sided
lower limit	-3.87
upper limit	-0.06
Variability estimate	Standard error of the mean
Dispersion value	0.961

Notes:

[1] - ANCOVA by dose using NRS baseline value and co-medication as covariates, with CFB to Week 12 (LOCF) as dependent variable was used to derive p-value.

<b>Statistical analysis title</b>	ANCOVA Analysis
Comparison groups	Placebo v MEDI7352 Medium Dose
Number of subjects included in analysis	68
Analysis specification	Pre-specified
Analysis type	other
P-value	= 0.1878 <sup>[2]</sup>
Method	ANCOVA
Parameter estimate	LS mean estimate
Point estimate	0.72
Confidence interval	
level	95 %
sides	2-sided
lower limit	-0.36
upper limit	1.79
Variability estimate	Standard error of the mean
Dispersion value	0.541

Notes:

[2] - ANCOVA by dose using NRS baseline value and co-medication as covariates, with CFB to Week 12 (LOCF) as dependent variable was used to derive p-value.

<b>Statistical analysis title</b>	ANCOVA Analysis
Comparison groups	Placebo v MEDI7352 High Dose
Number of subjects included in analysis	89
Analysis specification	Pre-specified
Analysis type	other
P-value	= 0.0009 <sup>[3]</sup>
Method	ANCOVA
Parameter estimate	LS mean estimate
Point estimate	-1.39
Confidence interval	
level	95 %
sides	2-sided
lower limit	-2.19
upper limit	-0.58

Variability estimate	Standard error of the mean
Dispersion value	0.407

Notes:

[3] - ANCOVA by dose using NRS baseline value and co-medication as covariates, with CFB to Week 12 (LOCF) as dependent variable was used to derive p-value.

### Secondary: Change From Baseline in Weekly Average of Average Daily Pain Score

End point title	Change From Baseline in Weekly Average of Average Daily Pain Score
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End point description:

Change from baseline to Weeks 2, 4, 6, 8, 10, and 18 in weekly average of average daily pain score is reported. Participants assessed their perceived daily average neuropathic pain over the previous 24 hours using an 11-point NRS, with 0 representing no pain and 10 representing the worst pain imaginable. Participants were instructed to assess their average daily pain at approximately the same time every morning, and to record the response in a subject diary (ePRO). mITT population included all participants who received at least 1 dose of any double-blind study drug and have at least 1 daily NRS assessment while receiving the treatment. Here, number analyzed (n) denotes those participants who were evaluable at the specified time point.

End point type	Secondary
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End point timeframe:

Baseline (Day -7 to Day -1, inclusive), Weeks 2, 4, 6, 8, 10, and 18

End point values	Placebo	MEDI7352 Low Dose	MEDI7352 Medium Dose	MEDI7352 High Dose
Subject group type	Reporting group	Reporting group	Reporting group	Reporting group
Number of subjects analysed	54	4	14	35
Units: Units on a scale				
arithmetic mean (standard deviation)				
Week 2 (n=53,4,14,35)	-0.385 (± 0.6863)	-1.089 (± 1.0258)	-0.311 (± 0.6285)	-0.540 (± 1.2503)
Week 4 (n=54,4,13,35)	-0.563 (± 1.0173)	-2.173 (± 1.4266)	-0.144 (± 0.9143)	-1.508 (± 1.5130)
Week 6 (n=50,4,13,34)	-0.955 (± 1.3236)	-2.458 (± 2.2156)	-0.578 (± 1.0715)	-2.044 (± 1.7916)
Week 8 (n=46,4,12,33)	-1.236 (± 1.6026)	-2.393 (± 2.0249)	-0.690 (± 1.2819)	-2.372 (± 2.0607)
Week 10 (n=44,4,12,33)	-1.418 (± 1.6274)	-2.595 (± 2.2003)	-0.712 (± 1.2164)	-2.629 (± 1.9943)
Week 18 (n=40,4,12,30)	-1.764 (± 1.8415)	-2.607 (± 2.2020)	-0.550 (± 1.2333)	-2.559 (± 2.1241)

### Statistical analyses

No statistical analyses for this end point

### Secondary: Percentage of Participants With $\geq 30\%$ and $\geq 50\%$ Reductions in Weekly Average of Average Daily Pain Score

End point title	Percentage of Participants With $\geq 30\%$ and $\geq 50\%$ Reductions in Weekly Average of Average Daily Pain Score
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End point description:

Percentage of participants with  $\geq 30\%$  and  $\geq 50\%$  decrease in weekly average of average daily pain score from baseline is reported. Participants assessed their perceived daily average neuropathic pain

over the previous 24 hours using an 11-point NRS, with 0 representing no pain and 10 representing the worst pain imaginable. Participants were instructed to assess their average daily pain at approximately the same time every morning, and to record the response in a subject diary (ePRO). mITT population included all participants who received at least 1 dose of any double-blind study drug and have at least 1 daily NRS assessment while receiving the treatment. Here, number analyzed (n) denotes those participants who were evaluable at the specified time point.

End point type	Secondary
End point timeframe:	
Baseline (Day -7 to Day -1, inclusive), Weeks 4, 8, 12, and 18 (follow-up)	

End point values	Placebo	MEDI7352 Low Dose	MEDI7352 Medium Dose	MEDI7352 High Dose
Subject group type	Reporting group	Reporting group	Reporting group	Reporting group
Number of subjects analysed	54	4	13	35
Units: Percentage of participants				
number (not applicable)				
>=30%: Week 4 (n=54,4,13,35)	3.7	75.0	7.7	34.3
>=30%: Week 8 (n=46,4,12,33)	28.3	75.0	16.7	57.6
>=30%: Week 12 (n=43,4,12,33)	32.6	75.0	8.3	66.7
>=30%: Week 18 (n=40,4,12,30)	35.0	50.0	25.0	56.7
>=50%: Week 4 (n=54,4,13,35)	1.9	0	0	14.3
>=50%: Week 8 (n=46,4,12,33)	13.0	25.0	8.3	36.4
>=50%: Week 12 (n=43,4,12,33)	11.6	50.0	8.3	42.4
>=50%: Week 18 (n=40,4,12,30)	15.0	25.0	0	40.0

## Statistical analyses

No statistical analyses for this end point

## Secondary: Change From Baseline in Galer Neuropathic Pain Scale (NPS)

End point title	Change From Baseline in Galer Neuropathic Pain Scale (NPS)
End point description:	
Change from baseline to Weeks 4, 8, 12, and 18 in Galer NPS total score is reported. Galer NPS included 2 descriptors of pain, including intensity and unpleasantness, and 8 descriptors that assessed specific qualities of neuropathic pain: sharp, hot, dull, cold, sensitive, itchy, deep, and surface pain. Each of these 10 dimensions had a 0 to 10 NRS in which 0 means no pain and 10 means most intense pain. Galer NPS total score (ranges from 0 to 100; with 0 and 100 representing no and highest degree of neuropathic-like symptoms, respectively) is sum of pain intensity, pain unpleasantness, pain sharpness, pain hotness, pain dullness, pain coldness, pain sensitivity, pain itching, deep pain intensity, and surface pain intensity (all in 11-point NRS). mITT population: participants who received at least 1 dose of any double-blind study drug and have at least 1 daily NRS assessment while receiving treatment. Number analyzed (n): participants who were evaluable at the specified time point.	
End point type	Secondary
End point timeframe:	
Baseline (Day -7 to Day -1, inclusive), Weeks 4, 8, 12, and 18 (follow-up)	

End point values	Placebo	MEDI7352 Low Dose	MEDI7352 Medium Dose	MEDI7352 High Dose
Subject group type	Reporting group	Reporting group	Reporting group	Reporting group
Number of subjects analysed	48	4	12	31
Units: Units on a scale				
arithmetic mean (standard deviation)				
Week 4 (n=48,4,11,31)	-13.333 (± 29.0461)	-5.750 (± 10.5317)	-10.727 (± 21.3967)	-26.387 (± 26.9922)
Week 8 (n=44,4,12,31)	-19.136 (± 31.5577)	-8.750 (± 18.2094)	-19.333 (± 23.7538)	-29.516 (± 32.3140)
Week 12 (n=39,4,11,29)	-24.385 (± 31.1081)	-28.750 (± 16.6808)	-22.000 (± 28.9379)	-34.586 (± 31.2622)
Week 18 (follow-up;n=36,4,11,29)	-25.722 (± 33.6151)	-23.000 (± 23.3095)	-27.909 (± 30.3725)	-25.414 (± 25.8947)

## Statistical analyses

No statistical analyses for this end point

## Secondary: Change From Baseline in Daily Sleep Interference Scale (DSIS)

End point title	Change From Baseline in Daily Sleep Interference Scale (DSIS)
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End point description:

The DSIS measures sleep interference due to pain. It is a single item measure that is completed by participants once a day (on waking up). It has an 11-point response scale (0-10) that asked participants to "select the number that best describes how much your pain has interfered with your sleep during the past 24 hours". Responses vary from 0 (did not interfere with sleep) to 10 (completely interfered with sleep-unable to sleep due to pain). Change from baseline to Weeks 4, 8, 12, and 18 (follow-up) in DSIS is reported. mITT population included all participants who received at least 1 dose of any double-blind study drug and have at least 1 daily NRS assessment while receiving the treatment. Number analyzed (n) denotes those participants who were evaluable at the specified time point.

End point type	Secondary
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End point timeframe:

Baseline (Day -7 to Day -1, inclusive), Weeks 4, 8, 12, and 18 (follow-up)

End point values	Placebo	MEDI7352 Low Dose	MEDI7352 Medium Dose	MEDI7352 High Dose
Subject group type	Reporting group	Reporting group	Reporting group	Reporting group
Number of subjects analysed	52	4	13	35
Units: Units on a scale				
arithmetic mean (standard deviation)				
Week 4 (n=52,4,13,35)	-0.760 (± 1.1212)	-2.810 (± 2.6979)	-0.979 (± 1.7930)	-1.790 (± 1.5688)
Week 8 (n=46,4,12,33)	-1.417 (± 1.7654)	-3.143 (± 2.7430)	-1.127 (± 1.4907)	-2.373 (± 2.1368)
Week 12 (n=42,4,12,33)	-1.668 (± 2.1928)	-3.735 (± 2.7598)	-1.077 (± 1.4769)	-2.725 (± 2.2484)
Week 18 (follow-up;n=40,4,12,30)	-1.824 (± 2.2582)	-3.036 (± 2.6557)	-0.865 (± 1.2795)	-2.602 (± 2.4427)

## Statistical analyses

No statistical analyses for this end point

### Secondary: Percentage of Participants With 'Improved', 'Much Improved', or 'Very Much Improved' Status in Patient Global Impression of Change (PGIC)

End point title	Percentage of Participants With 'Improved', 'Much Improved', or 'Very Much Improved' Status in Patient Global Impression of Change (PGIC)
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End point description:

Participants rated their overall improvement in health status using the PGIC. The PGIC consisted of a 7-point scale where 1 = very much improved and 7 = very much worse. The participants were asked the following question: How would you rate your overall improvement with treatment during the clinical study?, where the response options included the following: Very Much Improved, Much Improved, Minimally Improved, No Change, Minimally Worse, Much Worse, and Very Much Worse. mITT population included all participants who received at least 1 dose of any double-blind study drug and have at least 1 daily NRS assessment while receiving the treatment. Number analyzed (n) denotes those participants who were evaluable at the specified time point.

End point type	Secondary
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End point timeframe:

Baseline (Day -7 to Day -1, inclusive), Weeks 4, 8, 12, and 18 (follow-up)

End point values	Placebo	MEDI7352 Low Dose	MEDI7352 Medium Dose	MEDI7352 High Dose
Subject group type	Reporting group	Reporting group	Reporting group	Reporting group
Number of subjects analysed	48	4	12	31
Units: Percentage of participants				
number (not applicable)				
Week 4: Improved (n=48,4,11,31)	50.0	50.0	45.5	41.9
Week 4: Much Improved (n=48,4,11,31)	2.1	25.0	27.3	29.0
Week 4: Very Much Improved (n=48,4,11,31)	2.1	0	0	6.5
Week 8: Improved (n=44,4,12,31)	36.4	25.0	58.3	41.9
Week 8: Much Improved (n=44,4,12,31)	31.8	50.0	41.7	38.7
Week 8: Very Much Improved (n=44,4,12,31)	2.3	0	0	6.5
Week 12: Improved (n=39,4,11,29)	48.7	0	54.5	44.8
Week 12: Much Improved (n=39,4,11,29)	28.2	75.0	36.4	41.4
Week 12: Very Much Improved (n=39,4,11,29)	5.1	25.0	0	10.3
Week 18: Improved (n=36,4,11,29)	36.1	25.0	27.3	31.0
Week 18: Much Improved (n=36,4,11,29)	25.0	50.0	54.5	41.4
Week 18: Very Much Improved (n=36,4,11,29)	11.1	25.0	9.1	13.8

## Statistical analyses

**Secondary: Change From Baseline in 36-item Short-Form Health Survey (SF-36)**

End point title	Change From Baseline in 36-item Short-Form Health Survey (SF-36)
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## End point description:

Change from baseline to Week 12 in SF-36 is reported. The SF-36 assesses 8 health concepts: 1) limitations in physical activities because of health problems (physical functioning); 2) limitations in social activities because of physical or emotional problems (social functioning); 3) limitations in usual role activities because of physical health problems (role physical); 4) bodily pain; 5) general mental health; 6) limitations in usual role activities because of emotional problems (role emotional); 7) vitality; and 8) general health perceptions. The items use Likert-type scales with either 5 or 6 points, or 2 or 3 points. Higher SF-36 scores indicate a better state of health. Score for a section is an average of the individual question scores, which are scaled 0-100 (100=highest level of functioning). mITT population: all participants who received at least 1 dose of any double-blind study drug and have at least 1 daily NRS assessment while receiving treatment.

End point type	Secondary
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## End point timeframe:

Baseline (Day -7 to Day -1, inclusive) and Week 12

End point values	Placebo	MEDI7352 Low Dose	MEDI7352 Medium Dose	MEDI7352 High Dose
Subject group type	Reporting group	Reporting group	Reporting group	Reporting group
Number of subjects analysed	39	4	12	29
Units: Units on a scale				
arithmetic mean (standard deviation)				
Physical Functioning	11.282 (± 17.0412)	21.248 (± 13.1526)	0.417 (± 19.7084)	12.759 (± 16.6141)
Role Physical	7.372 (± 18.0159)	28.125 (± 27.7169)	0.000 (± 37.3101)	6.681 (± 17.4338)
Bodily Pain	8.3 (± 20.51)	9.8 (± 22.75)	12.1 (± 23.11)	14.9 (± 17.30)
General Health	-2.6 (± 17.70)	-2.5 (± 5.00)	-5.3 (± 17.72)	0.0 (± 19.98)
Vitality	-0.160 (± 20.6039)	3.125 (± 16.5359)	-0.521 (± 17.7695)	-4.095 (± 24.0486)
Social Functioning	1.92 (± 19.772)	31.25 (± 23.936)	-1.04 (± 24.108)	9.05 (± 23.121)
Role Emotional	-1.923 (± 25.3248)	8.333 (± 31.9110)	-5.556 (± 34.8748)	-1.724 (± 19.9681)
Mental Health	-5.9 (± 21.82)	10.0 (± 20.41)	-1.3 (± 14.48)	-0.7 (± 18.41)

**Statistical analyses**

No statistical analyses for this end point

**Secondary: Percentage of Participants Taking any Rescue Medication**

End point title	Percentage of Participants Taking any Rescue Medication
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## End point description:

Percentage of participants taking any rescue medication are reported. Participants were asked to record all rescue medications they take for neuropathic pain in a paper diary. mITT population included all participants who received at least 1 dose of any double-blind study drug and have at least 1 daily NRS assessment while receiving the treatment.

End point type	Secondary
End point timeframe:	
Baseline (Day -7 to Day -1, inclusive) through Week 12	

End point values	Placebo	MEDI7352 Low Dose	MEDI7352 Medium Dose	MEDI7352 High Dose
Subject group type	Reporting group	Reporting group	Reporting group	Reporting group
Number of subjects analysed	54	4	14	35
Units: Percentage of participants				
number (not applicable)	13.0	0	14.3	8.6

### Statistical analyses

No statistical analyses for this end point

### Secondary: Number of Participants With Treatment Emergent Adverse Events (TEAEs) and Treatment Emergent Serious Adverse Events (TESAEs)

End point title	Number of Participants With Treatment Emergent Adverse Events (TEAEs) and Treatment Emergent Serious Adverse Events (TESAEs)
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End point description:

An adverse event (AE) is any untoward medical occurrence in a participant who received study drug without regard to possibility of causal relationship. A serious adverse event (SAE) is an AE resulting in any of the following outcomes or deemed significant for any other reason: death; initial or prolonged inpatient hospitalization; life threatening experience (immediate risk of dying); persistent or significant disability/incapacity; congenital anomaly. The TEAEs are defined as events present at baseline that worsened in intensity after administration of study drug or events absent at baseline that emerged after administration of study drug. Safety population included all participants who received at least 1 dose of any double-blind study drug and were analyzed according to the treatment they actually received.

End point type	Secondary
End point timeframe:	
Day 1 through 20.42 weeks (maximum observed duration)	

End point values	Placebo	MEDI7352 Low Dose	MEDI7352 Medium Dose	MEDI7352 High Dose
Subject group type	Reporting group	Reporting group	Reporting group	Reporting group
Number of subjects analysed	54	4	14	35
Units: Participants				
Any TEAEs	30	4	10	23
Any TESAEs	1	0	0	0

### Statistical analyses

No statistical analyses for this end point

**Secondary: Number of Participants With Abnormal Vital Signs Reported as TEAEs**

End point title	Number of Participants With Abnormal Vital Signs Reported as TEAEs
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End point description:

Number of participants with abnormal vital signs reported as TEAEs are reported. Abnormal vital signs are defined as any abnormal finding in the vital sign parameters (body temperature, diastolic blood pressure, systolic blood pressure, heart rate, and respiratory rate).

End point type	Secondary
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End point timeframe:

Day 1 through 20.42 weeks (maximum observed duration)

End point values	Placebo	MEDI7352 Low Dose	MEDI7352 Medium Dose	MEDI7352 High Dose
Subject group type	Reporting group	Reporting group	Reporting group	Reporting group
Number of subjects analysed	54	4	14	35
Units: Participants				
Blood pressure increased	0	0	0	1
Hypotension	0	0	0	1
Hypertension	1	0	0	0
Orthostatic hypertension	1	0	0	0
Orthostatic hypotension	3	0	0	0

**Statistical analyses**

No statistical analyses for this end point

**Secondary: Number of Participants With Clinically Significant Abnormal Electrocardiograms (ECGs)**

End point title	Number of Participants With Clinically Significant Abnormal Electrocardiograms (ECGs)
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End point description:

Number of participants with clinically significant abnormal ECGs are reported. Safety population included all participants who received at least 1 dose of any double-blind study drug and were analyzed according to the treatment they actually received. Here, number analyzed (n) denotes those participants who were evaluable at the specified time point.

End point type	Secondary
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End point timeframe:

Day 1 through 20.42 weeks (maximum observed duration)

End point values	Placebo	MEDI7352 Low Dose	MEDI7352 Medium Dose	MEDI7352 High Dose
Subject group type	Reporting group	Reporting group	Reporting group	Reporting group
Number of subjects analysed	54	4	14	35
Units: Participants				
Day 1 (n=54,4,14,35)	1	0	0	0
Week 4 (n=50,4,13,32)	0	0	1	0



Week 8 (n=45,4,12,32)	0	0	0	1
Week 12 (n=40,4,11,31)	1	0	0	1
Week 18 (follow-up;n=40,4,12,30)	0	0	1	0

## Statistical analyses

No statistical analyses for this end point

## Secondary: Number of Participants With Abnormal Clinical Laboratory Parameters Reported as TEAEs

End point title	Number of Participants With Abnormal Clinical Laboratory Parameters Reported as TEAEs
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End point description:

Number of participants with abnormal clinical laboratory parameters reported as TEAEs are reported. Abnormal clinical laboratory parameters defined as any abnormal finding during analysis of hematology, clinical chemistry, coagulation, and urinalysis. Safety population included all participants who received at least 1 dose of any double-blind study drug and were analyzed according to the treatment they actually received.

End point type	Secondary
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End point timeframe:

Day 1 through 20.42 weeks (maximum observed duration)

End point values	Placebo	MEDI7352 Low Dose	MEDI7352 Medium Dose	MEDI7352 High Dose
Subject group type	Reporting group	Reporting group	Reporting group	Reporting group
Number of subjects analysed	54	4	14	35
Units: Participants				
Alanine aminotransferase increased	0	0	2	0
Blood glucose decreased	0	0	1	1
Aspartate aminotransferase increased	0	0	1	0
Blood glucose increased	0	0	0	1
C-reactive protein increased	1	0	0	1
Coagulation test abnormal	0	0	0	1
Glomerular filtration rate decreased	0	0	0	1
White blood cells urine positive	0	0	0	1
Bacterial test positive	1	0	0	0
Blood bilirubin increased	1	0	0	0
Protein urine present	1	0	0	0
Urine analysis abnormal	1	0	0	0
Hypoglycaemia	1	0	1	1
Hypercholesterolaemia	0	0	0	1
Hyperglycaemia	1	0	1	0

## Statistical analyses

No statistical analyses for this end point

### Secondary: Number of Participants With Clinically Significant Findings in Physical Examination Reported as TEAEs

End point title	Number of Participants With Clinically Significant Findings in Physical Examination Reported as TEAEs
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End point description:

Number of participants with clinically significant findings in physical examination reported as TEAEs are reported. A complete physical examination (excluding the genitourinary examination, unless warranted) was performed. Safety population included all participants who received at least 1 dose of any double-blind study drug and were analyzed according to the treatment they actually received.

End point type	Secondary
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End point timeframe:

Day 1 through 20.42 weeks (maximum observed duration)

End point values	Placebo	MEDI7352 Low Dose	MEDI7352 Medium Dose	MEDI7352 High Dose
Subject group type	Reporting group	Reporting group	Reporting group	Reporting group
Number of subjects analysed	54	4	14	35
Units: Participants	1	0	0	1

### Statistical analyses

No statistical analyses for this end point

### Secondary: Number of Participants With Clinically Significant Findings in Neurological Examination

End point title	Number of Participants With Clinically Significant Findings in Neurological Examination
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End point description:

Number of participants with clinically significant findings in neurological examination is reported. Safety population included all participants who received at least 1 dose of any double-blind study drug and were analyzed according to the treatment they actually received.

End point type	Secondary
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End point timeframe:

Day 1 through 20.42 weeks (maximum observed duration)

End point values	Placebo	MEDI7352 Low Dose	MEDI7352 Medium Dose	MEDI7352 High Dose
Subject group type	Reporting group	Reporting group	Reporting group	Reporting group
Number of subjects analysed	54	4	14	35
Units: Participants	0	0	0	0

## Statistical analyses

No statistical analyses for this end point

### Secondary: Number of Participants With Abnormal Dorsiflexion Strength

End point title	Number of Participants With Abnormal Dorsiflexion Strength
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End point description:

Number of participants with abnormal dorsiflexion strength is reported. Dorsiflexion strength is scored on 0-4 scale. The scale indicated 0 = normal power, 1 = mild weakness, 2 = moderate weakness, 3 = severe weakness, and 4 = paralysis. Safety population included all participants who received at least 1 dose of any double-blind study drug and were analyzed according to the treatment they actually received. Here, number of participants analyzed (N) denotes those participants who were assessed for strength and deep tendon reflexes assessment.

End point type	Secondary
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End point timeframe:

Day 1 through 20.42 weeks (maximum observed duration)

End point values	Placebo	MEDI7352 Low Dose	MEDI7352 Medium Dose	MEDI7352 High Dose
Subject group type	Reporting group	Reporting group	Reporting group	Reporting group
Number of subjects analysed	40	4	12	30
Units: Participants				
Mild Weakness	9	0	7	18
Moderate Weakness	8	0	0	2
Severe Weakness	0	0	0	0
Paralysis	0	0	0	0

## Statistical analyses

No statistical analyses for this end point

### Secondary: Number of Participants With Abnormal Deep Tendon Reflex (Knee and Ankle)

End point title	Number of Participants With Abnormal Deep Tendon Reflex (Knee and Ankle)
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End point description:

Number of participants with abnormal deep tendon reflex are reported. Deep tendon reflex (knee and ankle) strength is scored on 0-4 scale. The scale indicated 0 = normal, 1 = ankle reflex reduced, 2 = ankle reflex absent, 3 = ankle reflex absent and knee reflex reduced, and 4 = all reflexes (both ankle and knee) absent. Participants within a specific row are not included in any other row in the data table below. Safety population included all participants who received at least 1 dose of any double-blind study drug and were analyzed according to the treatment they actually received. Here, number of participants analyzed (N) denotes those participants who were assessed for strength and deep tendon reflexes assessment.

End point type	Secondary
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End point timeframe:

Day 1 through 20.42 weeks (maximum observed duration)

End point values	Placebo	MEDI7352 Low Dose	MEDI7352 Medium Dose	MEDI7352 High Dose
Subject group type	Reporting group	Reporting group	Reporting group	Reporting group
Number of subjects analysed	40	4	12	30
Units: Participants				
Ankle reflex reduced	5	2	2	3
Ankle reflex absent	4	0	1	5
Ankle reflex absent and knee reflex reduced	11	1	3	13
All reflexes absent	15	0	6	5

## Statistical analyses

No statistical analyses for this end point

## Secondary: Change From Baseline in Total Neuropathy Score-Nurse (TNSn)

End point title	Change From Baseline in Total Neuropathy Score-Nurse (TNSn)
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End point description:

TNSn, a semi-quantitative clinical assessment of peripheral nervous system function, was administered at baseline, Day 1, Weeks 2, 4, 6, 8, 10, 12, 18/early termination. It provides for assessment of motor and autonomic symptom, pin sensibility, and vibration sensibility score. Each neuropathy item is scored on 0–4 scale. The scores are summed to obtain a total score ranging 0–20, where score of 20 correlate with more severe neuropathy. Arbitrary numbers 99.999, 99999 signified mean and standard deviation (SD), respectively, not reported as no participants were evaluable in arm. Arbitrary number 999.99 signified SD not reported as only 1 participant was evaluable for this arm. Safety population: participants who received at least 1 dose of any double-blind study drug and were analyzed according to treatment actually received. Number analyzed (n): participants who were evaluable at specified time point. Only 3 early terminated participants contributed data at Week 18 assessment.

End point type	Secondary
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End point timeframe:

Baseline (Day -45 to -1), pre-dose on Day 1, Weeks 2, 4, 6, 8, 10, 12, and 18/early termination

End point values	Placebo	MEDI7352 Low Dose	MEDI7352 Medium Dose	MEDI7352 High Dose
Subject group type	Reporting group	Reporting group	Reporting group	Reporting group
Number of subjects analysed	50	4	13	34
Units: Units on a scale				
arithmetic mean (standard deviation)				
Day 1 (n=4,0,0,1)	-0.25 (± 0.957)	99.999 (± 99999)	99.999 (± 99999)	1.00 (± 999.99)
Week 2 (n=48,4,13,34)	-0.54 (± 1.304)	-1.25 (± 1.258)	0.00 (± 1.354)	-0.94 (± 2.117)
Week 4 (n=50,4,13,32)	-0.90 (± 1.982)	-2.50 (± 3.786)	-0.85 (± 1.819)	-1.19 (± 1.891)
Week 6 (n=45,4,12,33)	-1.40 (± 2.071)	-0.75 (± 2.872)	-1.00 (± 1.859)	-1.85 (± 2.093)
Week 8 (n=45,4,12,32)	-1.40 (± 2.082)	-2.50 (± 3.000)	-0.92 (± 1.881)	-1.84 (± 2.665)
Week 10 (n=40,4,12,31)	-1.90 (± 2.458)	-2.00 (± 3.916)	-1.67 (± 1.435)	-2.45 (± 3.042)
Week 12 (n=40,4,12,30)	-1.73 (± 2.562)	-2.25 (± 2.062)	-2.17 (± 1.899)	-2.53 (± 2.515)

Week 18 (n=40,4,12,30)	-2.00 (± 2.631)	-2.50 (± 3.512)	-2.17 (± 2.329)	-2.30 (± 3.053)
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## Statistical analyses

No statistical analyses for this end point

## Secondary: Number of Participants With Investigator Reported Significant Changes From Baseline in Motor and Sensory Nerve Conduction

End point title	Number of Participants With Investigator Reported Significant Changes From Baseline in Motor and Sensory Nerve Conduction
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End point description:

Motor and sensory nerve conduction studies were performed in relevant lower and upper limb nerves (sural, peroneal, median/ulnar, fibular, and tibial nerves) wherein amplitude, peak latency, conduction velocity, and duration of nerve action potentials were recorded. Safety population included all participants who received at least 1 dose of any double-blind study drug and were analyzed according to the treatment they actually received.

End point type	Secondary
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End point timeframe:

Day 1 through 20.42 weeks (maximum observed duration)

End point values	Placebo	MEDI7352 Low Dose	MEDI7352 Medium Dose	MEDI7352 High Dose
Subject group type	Reporting group	Reporting group	Reporting group	Reporting group
Number of subjects analysed	54	4	14	35
Units: Participants	2	0	0	0

## Statistical analyses

No statistical analyses for this end point

## Secondary: Number of Participants With at Least One Concomitant Medication

End point title	Number of Participants With at Least One Concomitant Medication
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End point description:

Number of participants with at least one concomitant medication is reported. A concomitant medication is defined as any medication continuing or starting after first dose of study medication. Safety population included all participants who received at least 1 dose of any double-blind study drug and were analyzed according to the treatment they actually received.

End point type	Secondary
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End point timeframe:

Day 1 through 20.42 weeks (maximum observed duration)

End point values	Placebo	MEDI7352 Low Dose	MEDI7352 Medium Dose	MEDI7352 High Dose
Subject group type	Reporting group	Reporting group	Reporting group	Reporting group
Number of subjects analysed	54	4	14	35
Units: Participants	54	4	14	35

## Statistical analyses

No statistical analyses for this end point

### Secondary: Number of Participants With Injection Site Reaction

End point title	Number of Participants With Injection Site Reaction
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End point description:

Number of participants with injection site reaction is reported. Safety population included all participants who received at least 1 dose of any double-blind study drug and were analyzed according to the treatment they actually received.

End point type	Secondary
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End point timeframe:

Day 1 through 20.42 weeks (maximum observed duration)

End point values	Placebo	MEDI7352 Low Dose	MEDI7352 Medium Dose	MEDI7352 High Dose
Subject group type	Reporting group	Reporting group	Reporting group	Reporting group
Number of subjects analysed	54	4	14	35
Units: Participants	1	1	0	0

## Statistical analyses

No statistical analyses for this end point

### Secondary: Number of Participants With Infusion Reaction

End point title	Number of Participants With Infusion Reaction
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End point description:

Number of participants with infusion reaction is reported. Safety population included all participants who received at least 1 dose of any double-blind study drug and were analyzed according to the treatment they actually received.

End point type	Secondary
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End point timeframe:

Day 1 through 20.42 weeks (maximum observed duration)

End point values	Placebo	MEDI7352 Low Dose	MEDI7352 Medium Dose	MEDI7352 High Dose
Subject group type	Reporting group	Reporting group	Reporting group	Reporting group
Number of subjects analysed	54	4	14	35
Units: Participants	0	0	0	2

## Statistical analyses

No statistical analyses for this end point

## Secondary: Serum Total Nerve Growth Factor (NGF) Concentrations

End point title	Serum Total Nerve Growth Factor (NGF) Concentrations
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End point description:

Serum concentrations of total NGF in ADA positive or negative participants are reported. Data for MEDI7352 low and medium dose group is not reported due to change in assay during the course of the study and stability data not supporting the re-analysis of early study samples with the new in-house assay. Safety population included all participants who received at least 1 dose of any double-blind study drug and were analyzed according to the treatment they actually received. Number of participants analyzed (N) denotes the number of participants evaluated for this outcome measure. Here, number analyzed (n) denotes those participants who were evaluable at the specified time point.

End point type	Secondary
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End point timeframe:

Pre-dose: Day 1 (baseline), Weeks 2, 4, 6, 8; Week 10 (Day 70; pre-dose, immediately before end of infusion, 8 hours post infusion start), Day 71 (post 24 hours of Day 70 infusion start), Weeks 11, 12 (approximately same time of day as Week 10 infusion)

End point values	Placebo	MEDI7352 Low Dose	MEDI7352 Medium Dose	MEDI7352 High Dose
Subject group type	Reporting group	Reporting group	Reporting group	Reporting group
Number of subjects analysed	30	0 <sup>[4]</sup>	0 <sup>[5]</sup>	31
Units: pg/mL				
arithmetic mean (standard deviation)				
Day 1 (n=30,0,0,31)	52.303 (± 11.6178)	()	()	65.687 (± 60.4036)
Week 2 (n=29,0,0,29)	62.613 (± 15.1088)	()	()	2384.760 (± 1989.0309)
Week 4 (n=29,0,0,29)	78.209 (± 92.0122)	()	()	2213.699 (± 2408.3112)
Week 6 (n=27,0,0,29)	64.661 (± 18.6478)	()	()	2361.855 (± 2645.8141)
Week 8 (n=26,0,0,29)	63.291 (± 16.5332)	()	()	1992.129 (± 2507.9826)
Week 10 (pre-dose; n=25,0,0,28)	173.383 (± 563.8921)	()	()	1681.073 (± 2455.0666)
Week 10 (before end of infusion; n=25,0,0,28)	57.308 (± 13.3588)	()	()	1992.756 (± 2968.1285)
Week 10 (8 hours post-dose; n=25,0,0,28)	177.323 (± 554.8131)	()	()	2012.384 (± 2795.8138)
Week 10 (post 24 hours; n=24,0,0,28)	222.980 (± 772.9408)	()	()	1849.835 (± 2671.9526)
Week 11 (n=25,0,0,26)	59.441 (± 13.6892)	()	()	1743.100 (± 2621.0687)

Week 12 (n=25,0,0,28)	63.416 ( $\pm$ 18.4243)	()	()	1710.966 ( $\pm$ 2838.0975)
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Notes:

[4] - Due to change in total NGF assay during study course, participants from this arm are not analysed.

[5] - Due to change in total NGF assay during study course, participants from this arm are not analysed.

## Statistical analyses

No statistical analyses for this end point

## Secondary: Number of Participants With Positive Anti-Drug Antibodies (ADA) to MEDI7352

End point title	Number of Participants With Positive Anti-Drug Antibodies (ADA) to MEDI7352
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End point description:

Number of participants with positive ADA to MEDI7352 are reported. Treatment-induced ADA positive is defined as ADA negative at baseline and positive at least 1 post-baseline assessment. Treatment-boosted ADA positive is defined as ADA positive at baseline with pre-existing titer boosted by 4-fold or greater during the study period. Persistent positive is defined as ADA negative at baseline and positive at least 2 post-baseline assessments (with  $\geq$  16 weeks between first and last positive) or ADA positive at last post-baseline assessment. Transiently positive is defined as ADA negative at baseline and at least one post-baseline ADA positive measurement and not fulfilling the conditions for persistently positive. Safety population included all participants who received at least 1 dose of any double-blind study drug and were analyzed according to the treatment they actually received. Here, number of participants analyzed (N) denotes those participants who had adequate ADA sample.

End point type	Secondary
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End point timeframe:

Pre-dose at baseline (Day -45 to -1) and Study Weeks 2, 4, 8, 10, 12, and 18 (follow-up)

End point values	Placebo	MEDI7352 Low Dose	MEDI7352 Medium Dose	MEDI7352 High Dose
Subject group type	Reporting group	Reporting group	Reporting group	Reporting group
Number of subjects analysed	53	4	14	35
Units: Participants				
Treatment-induced ADA	0	3	7	28
Treatment-boosted ADA	0	0	1	1
Persistently positive ADA	0	3	6	26
Transiently positive ADA	0	0	1	2

## Statistical analyses

No statistical analyses for this end point



## Adverse events

### Adverse events information

Timeframe for reporting adverse events:

Day 1 through 20.42 weeks (maximum observed duration)

Adverse event reporting additional description:

Safety population included all participants who received at least 1 dose of any double-blind study drug and were analyzed according to the treatment they actually received.

Assessment type	Systematic
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### Dictionary used

Dictionary name	MedDRA
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Dictionary version	26.0
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### Reporting groups

Reporting group title	Placebo
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Reporting group description:

Participants received intravenous (IV) placebo infusion matched to MEDI7352 during 12-week treatment period.

Reporting group title	MEDI7352 Medium Dose
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Reporting group description:

Participants received IV MEDI7352 during 12-week treatment period.

Reporting group title	MEDI7352 High Dose
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Reporting group description:

Participants received IV MEDI7352 during 12-week treatment period.

Reporting group title	MEDI7352 Low Dose
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Reporting group description:

Participants received IV MEDI7352 during 12-week treatment period.

Serious adverse events	Placebo	MEDI7352 Medium Dose	MEDI7352 High Dose
Total subjects affected by serious adverse events			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
number of deaths (all causes)	0	0	0
number of deaths resulting from adverse events	0	0	0
Infections and infestations			
Lung abscess			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0

Serious adverse events	MEDI7352 Low Dose		
Total subjects affected by serious adverse events			
subjects affected / exposed	0 / 4 (0.00%)		
number of deaths (all causes)	0		
number of deaths resulting from	0		

adverse events			
Infections and infestations			
Lung abscess			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences causally related to treatment / all	0 / 0		
deaths causally related to treatment / all	0 / 0		

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

<b>Non-serious adverse events</b>	Placebo	MEDI7352 Medium Dose	MEDI7352 High Dose
Total subjects affected by non-serious adverse events			
subjects affected / exposed	30 / 54 (55.56%)	10 / 14 (71.43%)	23 / 35 (65.71%)
Neoplasms benign, malignant and unspecified (incl cysts and polyps)			
Skin papilloma			
subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	1 / 35 (2.86%)
occurrences (all)	0	0	1
Vascular disorders			
Haematoma			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Lymphoedema			
subjects affected / exposed	0 / 54 (0.00%)	1 / 14 (7.14%)	0 / 35 (0.00%)
occurrences (all)	0	2	0
Hypotension			
subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	1 / 35 (2.86%)
occurrences (all)	0	0	1
Hypertension			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Orthostatic hypertension			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Orthostatic hypotension			
subjects affected / exposed	3 / 54 (5.56%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	3	0	0
General disorders and administration site conditions			

Feeling hot			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Chest pain			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Malaise			
subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	1 / 35 (2.86%)
occurrences (all)	0	0	1
Injection site reaction			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Gait disturbance			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Catheter site swelling			
subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	0	0	0
Facial pain			
subjects affected / exposed	0 / 54 (0.00%)	1 / 14 (7.14%)	0 / 35 (0.00%)
occurrences (all)	0	1	0
Gravitational oedema			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Respiratory, thoracic and mediastinal disorders			
Oropharyngeal pain			
subjects affected / exposed	1 / 54 (1.85%)	1 / 14 (7.14%)	1 / 35 (2.86%)
occurrences (all)	1	1	1
Rhinorrhoea			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Psychiatric disorders			
Bruxism			
subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	0	0	0
Insomnia			

subjects affected / exposed occurrences (all)	1 / 54 (1.85%) 1	0 / 14 (0.00%) 0	0 / 35 (0.00%) 0
Investigations			
Blood glucose increased subjects affected / exposed occurrences (all)	0 / 54 (0.00%) 0	0 / 14 (0.00%) 0	1 / 35 (2.86%) 1
Aspartate aminotransferase increased subjects affected / exposed occurrences (all)	0 / 54 (0.00%) 0	1 / 14 (7.14%) 1	0 / 35 (0.00%) 0
Blood glucose decreased subjects affected / exposed occurrences (all)	0 / 54 (0.00%) 0	1 / 14 (7.14%) 1	1 / 35 (2.86%) 1
Alanine aminotransferase increased subjects affected / exposed occurrences (all)	0 / 54 (0.00%) 0	2 / 14 (14.29%) 2	0 / 35 (0.00%) 0
Urine analysis abnormal subjects affected / exposed occurrences (all)	1 / 54 (1.85%) 1	0 / 14 (0.00%) 0	0 / 35 (0.00%) 0
Protein urine present subjects affected / exposed occurrences (all)	1 / 54 (1.85%) 1	0 / 14 (0.00%) 0	0 / 35 (0.00%) 0
Lymph node palpable subjects affected / exposed occurrences (all)	1 / 54 (1.85%) 1	0 / 14 (0.00%) 0	0 / 35 (0.00%) 0
Blood bilirubin increased subjects affected / exposed occurrences (all)	1 / 54 (1.85%) 1	0 / 14 (0.00%) 0	0 / 35 (0.00%) 0
Blood pressure increased subjects affected / exposed occurrences (all)	0 / 54 (0.00%) 0	0 / 14 (0.00%) 0	1 / 35 (2.86%) 1
White blood cells urine positive subjects affected / exposed occurrences (all)	0 / 54 (0.00%) 0	0 / 14 (0.00%) 0	1 / 35 (2.86%) 1
Glomerular filtration rate decreased			

subjects affected / exposed occurrences (all)	0 / 54 (0.00%) 0	0 / 14 (0.00%) 0	1 / 35 (2.86%) 1
Coagulation test abnormal subjects affected / exposed occurrences (all)	0 / 54 (0.00%) 0	0 / 14 (0.00%) 0	1 / 35 (2.86%) 1
C-reactive protein increased subjects affected / exposed occurrences (all)	1 / 54 (1.85%) 1	0 / 14 (0.00%) 0	1 / 35 (2.86%) 1
Bacterial test positive subjects affected / exposed occurrences (all)	1 / 54 (1.85%) 1	0 / 14 (0.00%) 0	0 / 35 (0.00%) 0
Injury, poisoning and procedural complications			
Repetitive strain injury subjects affected / exposed occurrences (all)	0 / 54 (0.00%) 0	0 / 14 (0.00%) 0	1 / 35 (2.86%) 1
Nail injury subjects affected / exposed occurrences (all)	0 / 54 (0.00%) 0	1 / 14 (7.14%) 1	0 / 35 (0.00%) 0
Infusion related reaction subjects affected / exposed occurrences (all)	0 / 54 (0.00%) 0	0 / 14 (0.00%) 0	2 / 35 (5.71%) 3
Contusion subjects affected / exposed occurrences (all)	1 / 54 (1.85%) 1	0 / 14 (0.00%) 0	0 / 35 (0.00%) 0
Fall subjects affected / exposed occurrences (all)	3 / 54 (5.56%) 3	0 / 14 (0.00%) 0	0 / 35 (0.00%) 0
Cardiac disorders			
Atrioventricular block first degree subjects affected / exposed occurrences (all)	1 / 54 (1.85%) 1	0 / 14 (0.00%) 0	0 / 35 (0.00%) 0
Ventricular extrasystoles subjects affected / exposed occurrences (all)	0 / 54 (0.00%) 0	0 / 14 (0.00%) 0	1 / 35 (2.86%) 1
Sinus tachycardia			

subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	1 / 35 (2.86%)
occurrences (all)	0	0	1
Palpitations			
subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	1 / 35 (2.86%)
occurrences (all)	0	0	1
Atrial fibrillation			
subjects affected / exposed	1 / 54 (1.85%)	1 / 14 (7.14%)	0 / 35 (0.00%)
occurrences (all)	1	2	0
Nervous system disorders			
Headache			
subjects affected / exposed	5 / 54 (9.26%)	1 / 14 (7.14%)	1 / 35 (2.86%)
occurrences (all)	9	2	1
Allodynia			
subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	1 / 35 (2.86%)
occurrences (all)	0	0	1
Paraesthesia			
subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	1 / 35 (2.86%)
occurrences (all)	0	0	1
Burning sensation			
subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	1 / 35 (2.86%)
occurrences (all)	0	0	1
Diabetic neuropathy			
subjects affected / exposed	2 / 54 (3.70%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	2	0	0
Carpal tunnel syndrome			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Sciatica			
subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	1 / 35 (2.86%)
occurrences (all)	0	0	1
Orthostatic intolerance			
subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	1 / 35 (2.86%)
occurrences (all)	0	0	1
Hypoaesthesia			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	1 / 35 (2.86%)
occurrences (all)	2	0	1

Dizziness subjects affected / exposed occurrences (all)	2 / 54 (3.70%) 2	0 / 14 (0.00%) 0	1 / 35 (2.86%) 1
Sensory disturbance subjects affected / exposed occurrences (all)	1 / 54 (1.85%) 1	0 / 14 (0.00%) 0	0 / 35 (0.00%) 0
Eye disorders			
Retinal haemorrhage subjects affected / exposed occurrences (all)	1 / 54 (1.85%) 1	0 / 14 (0.00%) 0	0 / 35 (0.00%) 0
Eye pain subjects affected / exposed occurrences (all)	0 / 54 (0.00%) 0	0 / 14 (0.00%) 0	0 / 35 (0.00%) 0
Eye irritation subjects affected / exposed occurrences (all)	0 / 54 (0.00%) 0	0 / 14 (0.00%) 0	0 / 35 (0.00%) 0
Eye haemorrhage subjects affected / exposed occurrences (all)	0 / 54 (0.00%) 0	1 / 14 (7.14%) 1	0 / 35 (0.00%) 0
Conjunctival hyperaemia subjects affected / exposed occurrences (all)	0 / 54 (0.00%) 0	1 / 14 (7.14%) 1	0 / 35 (0.00%) 0
Conjunctivitis allergic subjects affected / exposed occurrences (all)	0 / 54 (0.00%) 0	0 / 14 (0.00%) 0	0 / 35 (0.00%) 0
Gastrointestinal disorders			
Mouth ulceration subjects affected / exposed occurrences (all)	1 / 54 (1.85%) 1	1 / 14 (7.14%) 1	0 / 35 (0.00%) 0
Nausea subjects affected / exposed occurrences (all)	2 / 54 (3.70%) 2	1 / 14 (7.14%) 1	0 / 35 (0.00%) 0
Abdominal pain subjects affected / exposed occurrences (all)	1 / 54 (1.85%) 1	0 / 14 (0.00%) 0	0 / 35 (0.00%) 0
Change of bowel habit			

subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Frequent bowel movements			
subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	0	0	0
Food poisoning			
subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	1 / 35 (2.86%)
occurrences (all)	0	0	1
Diarrhoea			
subjects affected / exposed	5 / 54 (9.26%)	0 / 14 (0.00%)	1 / 35 (2.86%)
occurrences (all)	6	0	1
Constipation			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Abdominal discomfort			
subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	0	0	0
Dyspepsia			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Toothache			
subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	1 / 35 (2.86%)
occurrences (all)	0	0	1
Hypoaesthesia oral			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Melaena			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Abdominal pain upper			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Skin and subcutaneous tissue disorders			
Lichenification			
subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	1 / 35 (2.86%)
occurrences (all)	0	0	1



Hand dermatitis			
subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	1 / 35 (2.86%)
occurrences (all)	0	0	1
Eczema			
subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	1 / 35 (2.86%)
occurrences (all)	0	0	1
Dermatitis contact			
subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	1 / 35 (2.86%)
occurrences (all)	0	0	1
Dermatitis			
subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	1 / 35 (2.86%)
occurrences (all)	0	0	1
Erythema			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Skin burning sensation			
subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	1 / 35 (2.86%)
occurrences (all)	0	0	1
Pruritus			
subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	0	0	0
Renal and urinary disorders			
Urinary tract inflammation			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	1 / 35 (2.86%)
occurrences (all)	1	0	2
Musculoskeletal and connective tissue disorders			
Muscle spasms			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	2 / 35 (5.71%)
occurrences (all)	1	0	3
Muscle twitching			
subjects affected / exposed	0 / 54 (0.00%)	1 / 14 (7.14%)	0 / 35 (0.00%)
occurrences (all)	0	1	0
Pain in extremity			
subjects affected / exposed	2 / 54 (3.70%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	3	0	0
Soft tissue mass			

subjects affected / exposed	0 / 54 (0.00%)	1 / 14 (7.14%)	0 / 35 (0.00%)
occurrences (all)	0	1	0
Arthralgia			
subjects affected / exposed	4 / 54 (7.41%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	4	0	0
Back pain			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Musculoskeletal discomfort			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Musculoskeletal stiffness			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	2	0	0
Pain in jaw			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Infections and infestations			
Urinary tract infection			
subjects affected / exposed	2 / 54 (3.70%)	2 / 14 (14.29%)	1 / 35 (2.86%)
occurrences (all)	3	2	1
Upper respiratory tract infection			
subjects affected / exposed	3 / 54 (5.56%)	0 / 14 (0.00%)	4 / 35 (11.43%)
occurrences (all)	3	0	5
Herpes simplex			
subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	1 / 35 (2.86%)
occurrences (all)	0	0	1
Eye infection			
subjects affected / exposed	0 / 54 (0.00%)	1 / 14 (7.14%)	0 / 35 (0.00%)
occurrences (all)	0	1	0
Acute sinusitis			
subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	0	0	0
Nasopharyngitis			
subjects affected / exposed	11 / 54 (20.37%)	0 / 14 (0.00%)	2 / 35 (5.71%)
occurrences (all)	11	0	3

Viral upper respiratory tract infection			
subjects affected / exposed	0 / 54 (0.00%)	1 / 14 (7.14%)	0 / 35 (0.00%)
occurrences (all)	0	1	0
COVID-19			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Cellulitis			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Urinary tract infection bacterial			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Tooth abscess			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Respiratory tract infection			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Gastroenteritis			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Hordeolum			
subjects affected / exposed	1 / 54 (1.85%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	1	0	0
Metabolism and nutrition disorders			
Decreased appetite			
subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	0 / 35 (0.00%)
occurrences (all)	0	0	0
Hypoglycaemia			
subjects affected / exposed	1 / 54 (1.85%)	1 / 14 (7.14%)	1 / 35 (2.86%)
occurrences (all)	3	1	1
Hyperglycaemia			
subjects affected / exposed	1 / 54 (1.85%)	1 / 14 (7.14%)	0 / 35 (0.00%)
occurrences (all)	1	1	0
Hypercholesterolaemia			

subjects affected / exposed	0 / 54 (0.00%)	0 / 14 (0.00%)	1 / 35 (2.86%)
occurrences (all)	0	0	1

<b>Non-serious adverse events</b>	MEDI7352 Low Dose		
Total subjects affected by non-serious adverse events			
subjects affected / exposed	4 / 4 (100.00%)		
Neoplasms benign, malignant and unspecified (incl cysts and polyps)			
Skin papilloma			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Vascular disorders			
Haematoma			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Lymphoedema			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Hypotension			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Hypertension			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Orthostatic hypertension			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Orthostatic hypotension			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
General disorders and administration site conditions			
Feeling hot			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Chest pain			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		

Malaise subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		
Injection site reaction subjects affected / exposed occurrences (all)	1 / 4 (25.00%) 1		
Gait disturbance subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		
Catheter site swelling subjects affected / exposed occurrences (all)	1 / 4 (25.00%) 1		
Facial pain subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		
Gravitational oedema subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		
Respiratory, thoracic and mediastinal disorders Oropharyngeal pain subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		
Rhinorrhoea subjects affected / exposed occurrences (all)	1 / 4 (25.00%) 1		
Psychiatric disorders Bruxism subjects affected / exposed occurrences (all)	1 / 4 (25.00%) 1		
Insomnia subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		
Investigations Blood glucose increased subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		

Aspartate aminotransferase increased			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Blood glucose decreased			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Alanine aminotransferase increased			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Urine analysis abnormal			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Protein urine present			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Lymph node palpable			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Blood bilirubin increased			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Blood pressure increased			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
White blood cells urine positive			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Glomerular filtration rate decreased			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Coagulation test abnormal			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
C-reactive protein increased			

subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		
Bacterial test positive subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		
Injury, poisoning and procedural complications Repetitive strain injury subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		
Nail injury subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		
Infusion related reaction subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		
Contusion subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		
Fall subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		
Cardiac disorders Atrioventricular block first degree subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		
Ventricular extrasystoles subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		
Sinus tachycardia subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		
Palpitations subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		
Atrial fibrillation			

subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Nervous system disorders			
Headache			
subjects affected / exposed	3 / 4 (75.00%)		
occurrences (all)	24		
Allodynia			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Paraesthesia			
subjects affected / exposed	1 / 4 (25.00%)		
occurrences (all)	4		
Burning sensation			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Diabetic neuropathy			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Carpal tunnel syndrome			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Sciatica			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Orthostatic intolerance			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Hypoaesthesia			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Dizziness			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Sensory disturbance			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		



Eye disorders			
Retinal haemorrhage			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Eye pain			
subjects affected / exposed	1 / 4 (25.00%)		
occurrences (all)	1		
Eye irritation			
subjects affected / exposed	1 / 4 (25.00%)		
occurrences (all)	1		
Eye haemorrhage			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Conjunctival hyperaemia			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Conjunctivitis allergic			
subjects affected / exposed	1 / 4 (25.00%)		
occurrences (all)	1		
Gastrointestinal disorders			
Mouth ulceration			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Nausea			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Abdominal pain			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Change of bowel habit			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Frequent bowel movements			
subjects affected / exposed	1 / 4 (25.00%)		
occurrences (all)	1		
Food poisoning			

subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Diarrhoea			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Constipation			
subjects affected / exposed	1 / 4 (25.00%)		
occurrences (all)	1		
Abdominal discomfort			
subjects affected / exposed	1 / 4 (25.00%)		
occurrences (all)	1		
Dyspepsia			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Toothache			
subjects affected / exposed	1 / 4 (25.00%)		
occurrences (all)	1		
Hypoaesthesia oral			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Melaena			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Abdominal pain upper			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Skin and subcutaneous tissue disorders			
Lichenification			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Hand dermatitis			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Eczema			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		

Dermatitis contact subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		
Dermatitis subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		
Erythema subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		
Skin burning sensation subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		
Pruritus subjects affected / exposed occurrences (all)	1 / 4 (25.00%) 2		
Renal and urinary disorders Urinary tract inflammation subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		
Musculoskeletal and connective tissue disorders Muscle spasms subjects affected / exposed occurrences (all)	1 / 4 (25.00%) 1		
Muscle twitching subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		
Pain in extremity subjects affected / exposed occurrences (all)	1 / 4 (25.00%) 1		
Soft tissue mass subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		
Arthralgia subjects affected / exposed occurrences (all)	0 / 4 (0.00%) 0		
Back pain			

subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Musculoskeletal discomfort			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Musculoskeletal stiffness			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Pain in jaw			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Infections and infestations			
Urinary tract infection			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Upper respiratory tract infection			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Herpes simplex			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Eye infection			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Acute sinusitis			
subjects affected / exposed	1 / 4 (25.00%)		
occurrences (all)	1		
Nasopharyngitis			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Viral upper respiratory tract infection			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
COVID-19			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		

Cellulitis			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Urinary tract infection bacterial			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Tooth abscess			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Respiratory tract infection			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Gastroenteritis			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Hordeolum			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Metabolism and nutrition disorders			
Decreased appetite			
subjects affected / exposed	1 / 4 (25.00%)		
occurrences (all)	1		
Hypoglycaemia			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Hyperglycaemia			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		
Hypercholesterolaemia			
subjects affected / exposed	0 / 4 (0.00%)		
occurrences (all)	0		

## More information

### Substantial protocol amendments (globally)

Were there any global substantial amendments to the protocol? Yes

Date	Amendment
29 May 2019	<p>Added assessments of strength (dorsiflexion) and deep tendon reflexes (knee and ankle). Changed terminology from "cohort" to "stage". Added contraception guidance as an appendix. Modified the exclusion criteria to clarify that bilateral ankle radiographs were required at screening only if the participant had clinically significant sensory impairment. Modified the exclusion criteria to exclude participants with hemoglobin A1C (HbA1C) greater than 8.5% (formerly 8.0%). Specified that QuantiFERON tuberculosis (TB) testing was to be performed for all participants. Added a provision that, if necessary, the randomisation transaction in interactive web response system (IWRS) could be performed on Day -1. To avoid undue pressure on the participants, removed language that instructed the investigator to make every reasonable attempt to keep participants in the study. Added option for the sponsor to provide sterile water for injection and sodium chloride 0.9% solution for infusion if the site was unable to supply these. Noted that investigational product (IP) kit/code numbers were not to be recorded in the electronic case report form (eCRF). Removed the requirement that smoking habits were to remain unchanged for the duration of the study. Clarified that screening and Day 1 blood pressure assessments taken to assess orthostasis were to be measured in a supine (not sitting) position prior to the standing assessments. Clarified that urine pregnancy tests were to be performed in all women who were not surgically sterile. Specified that the sleep biosensor was to be worn on the nondominant wrist. Revised the list of sleep endpoints that the biosensor was capable of recording and analysing. Removed references to a clinical monitoring plan. Removed "Medimmune" from the sponsor name. Corrected typographical errors.</p>
21 February 2020	<p>Modified exclusion criteria to clarify that participants with a documented or suspected history of excessive alcohol usage or recreational drug usage should be evaluated at screening using appropriate quantitative or semi-quantitative methodologies according to local standards of accepted medical or psychiatric practice. Modified all instances of physical examinations to include physical and neurological examinations. Clarified that if the infusion was slowed or interrupted, the total infusion time should not exceed 4 hours at room temperature. Clarified that the timing of ECGs on Day 1 was relative to the start of IP administration. Added C-reactive protein to the serum chemistry tests to be performed. Modified the modified intent-to-treat population to include all randomised participants who received at least 1 dose of double-blind study medication and had at least 1 post-baseline (BL) NRS assessment.</p>
01 October 2020	<p>Modified doses used in Stage 3 of the study to include 10, 50, 150 and 450 µg/mL. Clarified the timing windows, namely for all vital signs, pharmacokinetic (PK), pharmacodynamic (PD), exploratory soluble biomarker sampling. Updated Clinical Experience from the most recent data safety cut-off of 23 April 2020. Modified the description of when participants in Stage 3 were to be randomly assigned to treatment of MEDI7352 or placebo. Modified the study design figure to include 10, 50, 150, and 450 µg/mL dose levels. Clarified the New York Heart Association (NYHA) exclusion criteria. Updated the number of study sites. Updated the study duration timing. Clarified who was to be included as an early terminated participant. Removed the futility analysis and clarified the naming of the Interim Analysis Review Committee to review the data for the interim analysis.</p>

04 December 2020	Added a fourth stage to the study and updated the number of participants to be included in each stage. Modified doses used in Stage 2 to 4 of the study. Clarified the timing windows, namely for all vital signs, PK, PD, exploratory soluble biomarker sampling. Updated Clinical Experience from the most recent data safety cut-off of 23 April 2020. Modified the description about the timing of randomization before Stage 3. Modified the study design figure to include the appropriate dosing during the 4 stages of the study. Modified the sample size description and number of participants needed in the study. Updated the efficacy analysis to include a supplementary analysis using an adaptive MCP-Mod method. Updated when participants could be replaced to the third and fourth stages of the study. Clarified the NYHA exclusion criteria. Updated the number of study sites. Updated the study duration. Clarified who was to be defined as an early terminated participant.
26 October 2021	Updated clinical experience with data from the completed and ongoing clinical studies as of 01 June 2021 in line with the updated investigator's brochure (Edition 6, dated 13 October 2021). Added text to exclusion criteria #11 to indicate that bilateral radiographs taken to evaluate osteoarthritis severity and exclude neuropathic arthropathy should include feet as well as ankles. Modified the definition of hypertension in exclusion criteria #27. Modified the HbA1C limit in exclusion criteria #29, from 8.5% to 10.0%. Evaluation of a cartilage degradation marker, urine C-terminal cross-linked telopeptide of type II collagen, was added. Corresponding changes were made to the exploratory endpoint describing soluble and genomic biomarkers to include this marker, with additional changes to the schedule of events and to text describing procedures at the appropriate visits. Adverse events of special interest (AESI) were added to align AESIs across studies per participant safety recommendations.
13 April 2022	Broadened electrocardiogram (ECG) language to allow paper ECGs as an alternative to digital ECGs. Clarified that the principal investigator's assessment of each ECG was to be recorded in the eCRF as normal, abnormal and clinically significant, or abnormal and not clinically significant. Clarified that urine samples for CTX-II biomarkers should only be collected from participants who had a baseline sample collected at the screening visit. Clarified the details relating to the interim analysis after completion of stage 3. Provided additional details of vital signs capture, particularly with regard to the use of a supine or sitting position.

Notes:

## Interruptions (globally)

Were there any global interruptions to the trial? Yes

Date	Interruption	Restart date
26 March 2020	Study halted temporarily due to COVID-19 pandemic	11 January 2021

Notes:

## Limitations and caveats

Limitations of the trial such as small numbers of subjects analysed or technical problems leading to unreliable data.

Dose-response relationship outcome was not evaluated by MCP-MOD because of insufficient number of doses. Study was prematurely terminated due to longer enrolment time. PK data analysed using population PK method; reported as separate report.

Notes: