



## Clinical trial results:

**The efficacy and safety of erdosteine in the long-term therapy of chronic obstructive pulmonary disease (COPD). A 12-month, randomised, double-blind, placebo-controlled, parallel group, multicenter study.**

### Summary

EudraCT number	2008-008192-34
Trial protocol	IT GB CZ FR DK SK BE BG
Global end of trial date	03 February 2014

### Results information

Result version number	v3 (current)
This version publication date	09 February 2017
First version publication date	13 November 2016
Version creation reason	<ul style="list-style-type: none"><li>• New data added to full data set</li><li>Safety data revision/others data correction</li></ul>
Summary attachment (see zip file)	Study Synopsis (RESTORE SYNOPSIS.pdf)

### Trial information

#### Trial identification

Sponsor protocol code	ERD-01-08/EP
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#### Additional study identifiers

ISRCTN number	-
ClinicalTrials.gov id (NCT number)	NCT01032304
WHO universal trial number (UTN)	-

Notes:

### Sponsors

Sponsor organisation name	Edmond Pharma Srl
Sponsor organisation address	S.S. dei Giovi 131, Paderno Dugnano (Mi), Italy, 20037
Public contact	Pozzi Edoardo, Edmond Pharma, +39 029500461, edoardo.pozzi@recipharm.com
Scientific contact	Pozzi Edoardo, Edmond Pharma, +39 029500461, edoardo.pozzi@recipharm.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	27 July 2016
Is this the analysis of the primary completion data?	No
Global end of trial reached?	Yes
Global end of trial date	03 February 2014
Was the trial ended prematurely?	No

Notes:

## General information about the trial

Main objective of the trial:

The principal objective of the study is to evaluate if erdosteine is effective in reducing the risk of acute exacerbations in patients affected by moderate-to-severe COPD during a 12-month treatment period.

Protection of trial subjects:

The efficacy of erdosteine has been evaluated using the experimental drug as add-on to a regular maintenance therapy of COPD. Patients continued their standard treatment for COPD, established by the physician according to the best clinical practice and guidelines. Since patients, in any case, receive the appropriate COPD standard treatment, the administration of placebo to half of them do not posed particular ethical concerns, while being at the same time scientifically sounded in order to establish the additional benefits of adding erdosteine to regular maintenance COPD treatment.

Background therapy: -

Evidence for comparator: -

Actual start date of recruitment	02 November 2009
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	United Kingdom: 9
Country: Number of subjects enrolled	Belgium: 9
Country: Number of subjects enrolled	Bulgaria: 12
Country: Number of subjects enrolled	Czech Republic: 96
Country: Number of subjects enrolled	Denmark: 12
Country: Number of subjects enrolled	France: 4
Country: Number of subjects enrolled	Italy: 94
Country: Number of subjects enrolled	Romania: 119
Country: Number of subjects enrolled	Poland: 101
Country: Number of subjects enrolled	Slovakia: 11
Worldwide total number of subjects	467
EEA total number of subjects	467

Notes:

### Subjects enrolled per age group

In utero	0
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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)	224
From 65 to 84 years	243
85 years and over	0

## Subject disposition

### Recruitment

Recruitment details:

First patient in 02-11-2009; last patient in 15/02/2013 . Countries involved: Italy, United Kingdom, Denmark, Czech Republic, France, Poland, Romania, Bulgaria, Slovakia and Belgium.

### Pre-assignment

Screening details:

Outpatients of both sexes, aged between 40 and 80 years with diagnosis of COPD (Stage II and III according to GOLD 2007); with a stable therapeutic regimen for COPD for at least 8 weeks prior to inclusion, 61 screening failure were registered due to consent withdrawn or lost to visit 1

### Pre-assignment period milestones

Number of subjects started	467
Number of subjects completed	467

### Period 1

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

### Arms

Are arms mutually exclusive? Yes

**Arm title** active

Arm description:

erdosteine cps 300 mg BID

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

Dosage and administration details:

300 mg BID

**Arm title** placebo

Arm description:

placebo controlled

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

Dosage and administration details:

Oral 300 mg BID

<b>Number of subjects in period 1</b>	active	placebo
Started	228	239
Completed	203	213
Not completed	25	26
Adverse event, serious fatal	1	2
Consent withdrawn by subject	14	20
Adverse event, non-fatal	10	4

## Baseline characteristics

### Reporting groups

Reporting group title	overall period
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Reporting group description: -

<b>Reporting group values</b>	overall period	Total	
Number of subjects	467	467	
Age categorical			
patient aged between 40 and 80 years			
Units: Subjects			
randomised	467	467	
not recorded	0	0	
Age continuous			
Units: years			
median	64.8		
standard deviation	± 8.34	-	
Gender categorical			
Units: Subjects			
Female	122	122	
Male	345	345	
Population characteristic			
Units: Subjects			
COPD	467	467	

## End points

### End points reporting groups

Reporting group title	active
Reporting group description:	erdosteine cps 300 mg BID
Reporting group title	placebo
Reporting group description:	placebo controlled

### Primary: overall number of exacerbations

End point title	overall number of exacerbations
End point description:	
End point type	Primary
End point timeframe:	treatment period

End point values	active	placebo		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	228	239		
Units: exact number				
exacerbations	1	1		

### Statistical analyses

Statistical analysis title	overall exacerbations rate
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Statistical analysis description:

A predefined analysis of exacerbation frequency was performed using a Poisson regression model, with the covariates of treatment, age, sex, Body Mass Index, and FEV1 at the baseline to estimate the rate ratio. The natural logarithm of the duration, expressed as months in the study, was used as an offset variable to correct for differences in the time individuals spent under observation.

Comparison groups	active v placebo
Number of subjects included in analysis	467
Analysis specification	Pre-specified
Analysis type	superiority
P-value	< 0.05
Method	Wilcoxon (Mann-Whitney)
Parameter estimate	Poisson regression estimate

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Confidence interval

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sides

2-sided

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## Adverse events

### Adverse events information

Timeframe for reporting adverse events:

treatment duration 12 months

Assessment type	Systematic
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### Dictionary used

Dictionary name	MedDRA
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Dictionary version	17.1
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### Reporting groups

Reporting group title	erdosteine
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Reporting group description: -

Reporting group title	placebo
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Reporting group description: -

<b>Serious adverse events</b>	erdosteine	placebo	
Total subjects affected by serious adverse events			
subjects affected / exposed	41 / 228 (17.98%)	33 / 239 (13.81%)	
number of deaths (all causes)	1	2	
number of deaths resulting from adverse events			
Investigations			
laryngoscopy			
subjects affected / exposed	1 / 228 (0.44%)	0 / 239 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Injury, poisoning and procedural complications			
carbonon monoxide poisoning			
alternative dictionary used: MedDRA 17.1			
alternative assessment type: Non-systematic			
subjects affected / exposed	1 / 228 (0.44%)	0 / 239 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Vascular disorders			
transient cerebrovascular insufficiency			
subjects affected / exposed	1 / 228 (0.44%)	0 / 239 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	

Transient ischaemic attack			
subjects affected / exposed	1 / 228 (0.44%)	0 / 239 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Aneurysm			
subjects affected / exposed	1 / 228 (0.44%)	0 / 239 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 1	0 / 0	
Stroke in evolution			
subjects affected / exposed	0 / 228 (0.00%)	1 / 239 (0.42%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Venous thrombosis			
subjects affected / exposed	1 / 228 (0.44%)	0 / 239 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Cardiac disorders			
atrial fibrillation			
subjects affected / exposed	1 / 228 (0.44%)	0 / 239 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Coronary artery disease			
subjects affected / exposed	1 / 228 (0.44%)	0 / 239 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
heart failure			
subjects affected / exposed	2 / 228 (0.88%)	0 / 239 (0.00%)	
occurrences causally related to treatment / all	0 / 2	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Angina pectoris			
subjects affected / exposed	0 / 228 (0.00%)	1 / 239 (0.42%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Arrhythmia			

subjects affected / exposed	0 / 228 (0.00%)	1 / 239 (0.42%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
<b>Surgical and medical procedures</b>			
Aortic bypass			
subjects affected / exposed	0 / 228 (0.00%)	1 / 239 (0.42%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
<b>Nervous system disorders</b>			
Depression			
subjects affected / exposed	0 / 228 (0.00%)	2 / 239 (0.84%)	
occurrences causally related to treatment / all	0 / 0	0 / 2	
deaths causally related to treatment / all	0 / 0	0 / 0	
<b>Blood and lymphatic system disorders</b>			
Thrombocytopenia			
subjects affected / exposed	1 / 228 (0.44%)	0 / 239 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
<b>Gastrointestinal disorders</b>			
chest/epigastric pain			
subjects affected / exposed	1 / 228 (0.44%)	0 / 239 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
<b>Hepatobiliary disorders</b>			
Cholecystectomy			
subjects affected / exposed	1 / 228 (0.44%)	0 / 239 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Gallbladder empyema			
subjects affected / exposed	0 / 228 (0.00%)	1 / 239 (0.42%)	
occurrences causally related to treatment / all	0 / 0	1 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
<b>Respiratory, thoracic and mediastinal disorders</b>			
COPD exacerbation			

subjects affected / exposed	16 / 228 (7.02%)	16 / 239 (6.69%)	
occurrences causally related to treatment / all	0 / 22	0 / 18	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pneumonia			
subjects affected / exposed	4 / 228 (1.75%)	5 / 239 (2.09%)	
occurrences causally related to treatment / all	0 / 4	0 / 5	
deaths causally related to treatment / all	0 / 0	0 / 0	
Acute respiratory failure			
subjects affected / exposed	1 / 228 (0.44%)	1 / 239 (0.42%)	
occurrences causally related to treatment / all	0 / 1	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 1	
Pneumonia aspiration			
subjects affected / exposed	0 / 228 (0.00%)	1 / 239 (0.42%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 1	
Pneumothorax			
subjects affected / exposed	0 / 228 (0.00%)	1 / 239 (0.42%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Chest pain			
subjects affected / exposed	0 / 228 (0.00%)	1 / 239 (0.42%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pulmonary oedema			
subjects affected / exposed	0 / 228 (0.00%)	1 / 239 (0.42%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Renal and urinary disorders			
cancer			
subjects affected / exposed	1 / 228 (0.44%)	0 / 239 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Urinary tract infection			

subjects affected / exposed	1 / 228 (0.44%)	0 / 239 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Urinary retention			
subjects affected / exposed	0 / 228 (0.00%)	1 / 239 (0.42%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Musculoskeletal and connective tissue disorders			
vertebral pain			
subjects affected / exposed	1 / 228 (0.44%)	0 / 239 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Back pain			
subjects affected / exposed	0 / 228 (0.00%)	1 / 239 (0.42%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hip fracture			
subjects affected / exposed	0 / 228 (0.00%)	1 / 239 (0.42%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	

Frequency threshold for reporting non-serious adverse events: 1 %

<b>Non-serious adverse events</b>	erdosteine	placebo	
Total subjects affected by non-serious adverse events			
subjects affected / exposed	228 / 228 (100.00%)	206 / 239 (86.19%)	
Surgical and medical procedures			
Infiltration anaesthesia			
subjects affected / exposed	1 / 228 (0.44%)	0 / 239 (0.00%)	
occurrences (all)	1	0	
General disorders and administration site conditions			
Liver function test abnormal			

subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	2 / 239 (0.84%) 2	
Chills subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	0 / 239 (0.00%) 0	
Fatigue subjects affected / exposed occurrences (all)	2 / 228 (0.88%) 2	0 / 239 (0.00%) 0	
Asthenia subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	1 / 239 (0.42%) 1	
Haemoptysis subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	0 / 239 (0.00%) 0	
Headache subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	4 / 239 (1.67%) 5	
Insomnia subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	2 / 239 (0.84%) 2	
Tooth abscess subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 2	1 / 239 (0.42%) 1	
Toothache subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	1 / 239 (0.42%) 2	
Weight decreased subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	0 / 239 (0.00%) 0	
Respiratory, thoracic and mediastinal disorders			
Viral infection subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	0 / 239 (0.00%) 0	
Bronchopneumopathy			

subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	0 / 239 (0.00%) 0
Rhinitis		
subjects affected / exposed occurrences (all)	5 / 228 (2.19%) 5	4 / 239 (1.67%) 6
Chest pain		
subjects affected / exposed occurrences (all)	2 / 228 (0.88%) 2	0 / 239 (0.00%) 0
Cough		
subjects affected / exposed occurrences (all)	2 / 228 (0.88%) 2	3 / 239 (1.26%) 3
Throat irritation		
subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	4 / 239 (1.67%) 4
Dyspnoea		
subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	8 / 239 (3.35%) 9
Sputum discoloured		
subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	0 / 239 (0.00%) 0
Ear infection		
subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	1 / 239 (0.42%) 1
Painful respiration		
subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	0 / 239 (0.00%) 0
Sinusitis		
subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	1 / 239 (0.42%) 1
Upper respiratory tract infection		
subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	1 / 239 (0.42%) 1
COPD exacerbations		
subjects affected / exposed occurrences (all)	99 / 228 (43.42%) 185	136 / 239 (56.90%) 258
Investigations		

Mitral valve replacement subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	1 / 239 (0.42%) 1	
Injury, poisoning and procedural complications Chest injury subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	1 / 239 (0.42%) 1	
Cardiac disorders Atrial fibrillation subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	1 / 239 (0.42%) 1	
Cardiac disorder subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	0 / 239 (0.00%) 0	
Palpitations subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	2 / 239 (0.84%) 2	
Arrhythmia subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	1 / 239 (0.42%) 1	
Coronary artery disease subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	1 / 239 (0.42%) 1	
Hypertension subjects affected / exposed occurrences (all)	8 / 228 (3.51%) 9	2 / 239 (0.84%) 2	
Hypocalcaemia subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	1 / 239 (0.42%) 1	
Tachycardia paroxysmal subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	1 / 239 (0.42%) 1	
Nervous system disorders Anxiety subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	0 / 239 (0.00%) 0	

Depression subjects affected / exposed occurrences (all)	2 / 228 (0.88%) 3	0 / 239 (0.00%) 0	
Dizziness subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	1 / 239 (0.42%) 1	
Tremor subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	1 / 239 (0.42%) 1	
Sleep disorder subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	1 / 239 (0.42%) 1	
Blood and lymphatic system disorders			
Anaemia subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	0 / 239 (0.00%) 0	
Hypoxia subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	1 / 239 (0.42%) 1	
Arterial disorder subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 2	0 / 239 (0.00%) 0	
Eye disorders			
Cataract subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 2	0 / 239 (0.00%) 0	
Corneal abrasion subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	0 / 239 (0.00%) 0	
Pain subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	0 / 239 (0.00%) 0	
Gastrointestinal disorders			
Abdominal pain subjects affected / exposed occurrences (all)	2 / 228 (0.88%) 2	3 / 239 (1.26%) 3	
Reflux gastritis			

subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	1 / 239 (0.42%) 1	
Diarrhoea subjects affected / exposed occurrences (all)	2 / 228 (0.88%) 3	1 / 239 (0.42%) 1	
Dyspepsia subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	6 / 239 (2.51%) 6	
Gastric ulcer subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	0 / 239 (0.00%) 0	
Acidosis subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	1 / 239 (0.42%) 1	
Nausea subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	1 / 239 (0.42%) 1	
Vomiting subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	0 / 239 (0.00%) 0	
Hepatobiliary disorders Bladder spasm subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	0 / 239 (0.00%) 0	
Cholecystitis subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	1 / 239 (0.42%) 1	
Salivary hypersecretion subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	1 / 239 (0.42%) 1	
Skin and subcutaneous tissue disorders Pruritus subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	1 / 239 (0.42%) 1	
Rash			

subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	1 / 239 (0.42%) 1	
Infection subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	0 / 239 (0.00%) 0	
Renal and urinary disorders Urinary retention subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	0 / 239 (0.00%) 0	
Cystitis subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	1 / 239 (0.42%) 1	
Urinary tract inflammation subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	1 / 239 (0.42%) 1	
Proteinuria subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	1 / 239 (0.42%) 1	
Urinary tract infection subjects affected / exposed occurrences (all)	4 / 228 (1.75%) 4	1 / 239 (0.42%) 0	
Incontinence subjects affected / exposed occurrences (all)	4 / 228 (1.75%) 4	1 / 239 (0.42%) 1	
Endocrine disorders Diabetes mellitus subjects affected / exposed occurrences (all)	2 / 228 (0.88%) 2	1 / 239 (0.42%) 1	
Musculoskeletal and connective tissue disorders Back pain subjects affected / exposed occurrences (all)	3 / 228 (1.32%) 3	3 / 239 (1.26%) 3	
Contusion subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	1 / 239 (0.42%) 1	
Bite			

subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	1 / 239 (0.42%) 1	
Fibromyalgia subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	2 / 239 (0.84%) 2	
Joint abscess subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	1 / 239 (0.42%) 1	
Bone pain subjects affected / exposed occurrences (all)	2 / 228 (0.88%) 2	1 / 239 (0.42%) 1	
Oedema subjects affected / exposed occurrences (all)	2 / 228 (0.88%) 2	1 / 239 (0.42%) 1	
Infections and infestations			
Influenza subjects affected / exposed occurrences (all)	3 / 228 (1.32%) 3	3 / 239 (1.26%) 3	
Pyrexia subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	0 / 239 (0.00%) 0	
Mycobacterial infection subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	0 / 239 (0.00%) 0	
Fungal infection subjects affected / exposed occurrences (all)	0 / 228 (0.00%) 0	2 / 239 (0.84%) 2	
Pneumonia subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	1 / 239 (0.42%) 1	
Viraemia subjects affected / exposed occurrences (all)	1 / 228 (0.44%) 1	2 / 239 (0.84%) 2	
Metabolism and nutrition disorders Hyper HDL cholesterolaemia			

subjects affected / exposed	1 / 228 (0.44%)	0 / 239 (0.00%)	
occurrences (all)	1	0	
Cachexia			
subjects affected / exposed	0 / 228 (0.00%)	1 / 239 (0.42%)	
occurrences (all)	0	1	

## **More information**

### **Substantial protocol amendments (globally)**

Were there any global substantial amendments to the protocol? No

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### **Interruptions (globally)**

Were there any global interruptions to the trial? No

### **Limitations and caveats**

None reported