



## Clinical trial results: Post Marketing Surveillance Study to Observe the Safety and Efficacy of Voriconazole Intravenous Infusion

### Summary

EudraCT number	2017-000197-11
Trial protocol	Outside EU/EEA
Global end of trial date	15 May 2012

### Results information

Result version number	v1 (current)
This version publication date	08 June 2017
First version publication date	08 June 2017

### Trial information

#### Trial identification

Sponsor protocol code	A1501067
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#### Additional study identifiers

ISRCTN number	-
ClinicalTrials.gov id (NCT number)	NCT01073618
WHO universal trial number (UTN)	-

Notes:

### Sponsors

Sponsor organisation name	Pfizer, Inc.
Sponsor organisation address	235 E 42nd Street, New York, United States, 110017
Public contact	Pfizer ClinicalTrials.gov Call Center, Pfizer, Inc., 001 800 718 1021, ClinicalTrials.gov_Inquiries@pfizer.com
Scientific contact	Pfizer ClinicalTrials.gov Call Center, Pfizer, Inc., 001 18007181021, ClinicalTrials.gov_Inquiries@pfizer.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?	Yes

Notes:

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**Results analysis stage**

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Analysis stage	Final
Date of interim/final analysis	19 October 2009
Is this the analysis of the primary completion data?	No

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Global end of trial reached?	Yes
Global end of trial date	15 May 2012
Was the trial ended prematurely?	No

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Notes:

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**General information about the trial**

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Main objective of the trial:

The objective of this study was to determine the efficacy and safety of Voriconazole, 200 mg intravenous infusion (IV).

Protection of trial subjects:

The study was in compliance with the ethical principles derived from the Declaration of Helsinki and in compliance with all International Council for Harmonization (ICH) Good Clinical Practice (GCP) Guidelines. All the local regulatory requirements pertinent to safety of trial subjects were followed.

Background therapy: -

Evidence for comparator: -

Actual start date of recruitment	11 April 2006
Long term follow-up planned	Yes
Long term follow-up rationale	Safety, Efficacy
Long term follow-up duration	2 Years
Independent data monitoring committee (IDMC) involvement?	No

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Notes:

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**Population of trial subjects**

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**Subjects enrolled per country**

Country: Number of subjects enrolled	Korea, Republic of: 692
Worldwide total number of subjects	692
EEA total number of subjects	0

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Notes:

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**Subjects enrolled per age group**

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In utero	0
Preterm newborn - gestational age < 37 wk	0
Newborns (0-27 days)	0
Infants and toddlers (28 days-23 months)	10
Children (2-11 years)	45
Adolescents (12-17 years)	42
Adults (18-64 years)	500
From 65 to 84 years	95

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85 years and over	0
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## Subject disposition

### Recruitment

Recruitment details: -

### Pre-assignment

Screening details:

A total of 692 subjects were enrolled at multiple centers in Republic of Korea in this study. The study was conducted from 11-Apr-2006 to 19-Oct-2009.

### Period 1

Period 1 title	Overall Study (overall period)
Is this the baseline period?	Yes
Allocation method	Not applicable
Blinding used	Not blinded

### Arms

Arm title	Voriconazole
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Arm description:

Subjects aged greater than (>) 12 years with invasive aspergillosis received Voriconazole intravenously at a loading dose of 6 milligrams per kilogram (mg/kg), every 12 hours for the first 24 hours, followed by the maintenance dose of 4 mg/kg intravenously, twice daily up to 2 weeks. Subjects aged >12 years with invasive candidemia received Voriconazole intravenously at a loading dose of 6 mg/kg intravenously, every 12 hours for the first 24 hours, followed by the maintenance dose of Voriconazole 3 to 4 mg/kg intravenously, twice daily up to 2 weeks.

Arm type	Experimental
Investigational medicinal product name	Voriconazole 6 mg/kg
Investigational medicinal product code	Voriconazole
Other name	
Pharmaceutical forms	Powder for solution for injection/infusion
Routes of administration	Intravenous use

Dosage and administration details:

Subjects received loading dose of Voriconazole 6 mg/kg intravenously twice daily for the first 24 hours up to 2 weeks.

Investigational medicinal product name	Voriconazole 4 mg/kg
Investigational medicinal product code	Voriconazole
Other name	
Pharmaceutical forms	Powder for solution for injection/infusion
Routes of administration	Intravenous use

Dosage and administration details:

Subjects received maintenance dose of Voriconazole 4 mg/kg intravenously twice daily up to 2 weeks.

Number of subjects in period 1	Voriconazole
Started	692
Completed	385
Not completed	307
Adverse Event	19
Death	208
Unspecified	80



## Baseline characteristics

### Reporting groups

Reporting group title	Voriconazole
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Reporting group description:

Subjects aged greater than (>) 12 years with invasive aspergillosis received Voriconazole intravenously at a loading dose of 6 milligrams per kilogram (mg/kg), every 12 hours for the first 24 hours, followed by the maintenance dose of 4 mg/kg intravenously, twice daily up to 2 weeks. Subjects aged >12 years with invasive candidemia received Voriconazole intravenously at a loading dose of 6 mg/kg intravenously, every 12 hours for the first 24 hours, followed by the maintenance dose of Voriconazole 3 to 4 mg/kg intravenously, twice daily up to 2 weeks.

Reporting group values	Voriconazole	Total	
Number of subjects	692	692	
Age Categorical			
Units: Subjects			
<18 years	97	97	
Between 18 and 44 years	218	218	
Between 45 and 64 years	282	282	
>=65 years	95	95	
Age continuous			
Units: years			
arithmetic mean	43.7		
standard deviation	± 19.7	-	
Gender, Male/Female			
Units: Subjects			
Female	275	275	
Male	417	417	

## End points

### End points reporting groups

Reporting group title	Voriconazole
Reporting group description:	
Subjects aged greater than (>) 12 years with invasive aspergillosis received Voriconazole intravenously at a loading dose of 6 milligrams per kilogram (mg/kg), every 12 hours for the first 24 hours, followed by the maintenance dose of 4 mg/kg intravenously, twice daily up to 2 weeks. Subjects aged >12 years with invasive candidemia received Voriconazole intravenously at a loading dose of 6 mg/kg intravenously, every 12 hours for the first 24 hours, followed by the maintenance dose of Voriconazole 3 to 4 mg/kg intravenously, twice daily up to 2 weeks.	

### Primary: Percentage of Subjects With Categorical Clinical Response: Cure, Improvement, Failure, or Unevaluable

End point title	Percentage of Subjects With Categorical Clinical Response: Cure, Improvement, Failure, or Unevaluable <sup>[1]</sup>
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End point description:

Clinical response was defined as: Cure = resolution of all baseline signs and symptoms of fungal infection(s); Improvement = lessening of baseline signs and symptoms or absence of one or more, but not all baseline findings; Failure = no improvement or deterioration of baseline condition; Unevaluable = Incomplete therapy (efficacy could not be evaluated or discontinuation was not followed up). Intent to treat (ITT) population included subjects who received study drug for the approved indication and had been evaluated for related parameters at least once.

End point type	Primary
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End point timeframe:

Baseline (Day 1) up to 2 years

Notes:

[1] - No statistical analyses have been specified for this primary end point. It is expected there is at least one statistical analysis for each primary end point.

Justification: Only descriptive analysis was planned to be analyzed in this endpoint

End point values	Voriconazole			
Subject group type	Reporting group			
Number of subjects analysed	692			
Units: percentage of subjects				
number (not applicable)				
Cure	12.72			
Improvement	41.47			
Failure	17.92			
Unevaluable	27.89			

### Statistical analyses

No statistical analyses for this end point

### Secondary: Percentage of Subjects With Cultivated Strain Mycological Response: Eradication, Persistence, Superinfection, or Not evaluable

End point title	Percentage of Subjects With Cultivated Strain Mycological Response: Eradication, Persistence, Superinfection, or Not evaluable
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**End point description:**

Mycological response was defined as: Eradication = absence of signs and symptoms of fungal infection; Persistence = (no eradication) presence of fungal infection; Superinfection = existence of different strains from strains separated prior to study medication; Not evaluable = a follow-up mycological cultivation is not performed. ITT population included subjects who received study drug for the approved indication and had been evaluated for related parameters at least once. Here, 'N' (Number of subjects analyzed) signifies those subjects who were evaluable for this endpoint.

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End point type	Secondary
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End point timeframe:

Baseline (Day 1) up to 2 years

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End point values	Voriconazole			
Subject group type	Reporting group			
Number of subjects analysed	159			
Units: percentage of subjects				
number (not applicable)				
Eradication	64.78			
Persistence	27.67			
Superinfection	3.77			
Not evaluable	3.77			

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**Statistical analyses**

No statistical analyses for this end point

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## Adverse events

### Adverse events information

Timeframe for reporting adverse events:

Baseline up to 28 days after last dose of study drug

Adverse event reporting additional description:

The same event may appear as both an AE and a SAE. However, what is presented are distinct events. An event may be categorized as serious in one subject and as nonserious in another subject, or one subject may have experienced both a serious and nonserious event during the study.

Assessment type	Systematic
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### Dictionary used

Dictionary name	MedDRA
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Dictionary version	13.1
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### Reporting groups

Reporting group title	Voriconazole
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Reporting group description:

Subjects aged >12 years with invasive aspergillosis received Voriconazole intravenously at a loading dose of 6 mg/kg, every 12 hours for the first 24 hours, followed by the maintenance dose of 4 mg/kg intravenously, twice daily up to 2 weeks. Subjects aged >12 years with invasive candidemia received Voriconazole intravenously at a loading dose of 6 mg/kg intravenously, every 12 hours for the first 24 hours, followed by the maintenance dose of Voriconazole 3 to 4 mg/kg intravenously, twice daily up to 2 weeks.

Serious adverse events	Voriconazole		
Total subjects affected by serious adverse events			
subjects affected / exposed	239 / 692 (34.54%)		
number of deaths (all causes)	220		
number of deaths resulting from adverse events			
Neoplasms benign, malignant and unspecified (incl cysts and polyps)			
Acute lymphocytic leukaemia			
subjects affected / exposed	2 / 692 (0.29%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	1 / 1		
Acute myeloid leukaemia			
subjects affected / exposed	7 / 692 (1.01%)		
occurrences causally related to treatment / all	0 / 7		
deaths causally related to treatment / all	0 / 7		
Hypopharyngeal cancer			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	1 / 1		

Leukaemia				
subjects affected / exposed	18 / 692 (2.60%)			
occurrences causally related to treatment / all	0 / 18			
deaths causally related to treatment / all	0 / 18			
Leukaemia recurrent				
subjects affected / exposed	1 / 692 (0.14%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 1			
Leukaemic infiltration				
subjects affected / exposed	1 / 692 (0.14%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Lung neoplasm malignant				
subjects affected / exposed	1 / 692 (0.14%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 1			
Lymphoma				
subjects affected / exposed	1 / 692 (0.14%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 1			
Multiple myeloma				
subjects affected / exposed	1 / 692 (0.14%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Tumour haemorrhage				
subjects affected / exposed	1 / 692 (0.14%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 1			
Rectal cancer				
subjects affected / exposed	1 / 692 (0.14%)			
occurrences causally related to treatment / all	1 / 1			
deaths causally related to treatment / all	1 / 1			
Myelodysplastic syndrome				

subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
Vascular disorders			
Hypotension			
subjects affected / exposed	3 / 692 (0.43%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 1		
Shock			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
General disorders and administration site conditions			
Brain death			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
Condition aggravated			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
Disease progression			
subjects affected / exposed	14 / 692 (2.02%)		
occurrences causally related to treatment / all	3 / 14		
deaths causally related to treatment / all	3 / 14		
General physical health deterioration			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
Multi-organ failure			
subjects affected / exposed	17 / 692 (2.46%)		
occurrences causally related to treatment / all	0 / 17		
deaths causally related to treatment / all	0 / 14		

Pyrexia			
subjects affected / exposed	2 / 692 (0.29%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
Immune system disorders			
Acute graft versus host disease in intestine			
subjects affected / exposed	2 / 692 (0.29%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
Acute graft versus host disease in skin			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Graft versus host disease			
subjects affected / exposed	2 / 692 (0.29%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
Respiratory, thoracic and mediastinal disorders			
Acute respiratory distress syndrome			
subjects affected / exposed	12 / 692 (1.73%)		
occurrences causally related to treatment / all	1 / 12		
deaths causally related to treatment / all	1 / 9		
Acute respiratory failure			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
Bronchopulmonary dysplasia			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
Dyspnoea			

subjects affected / exposed	9 / 692 (1.30%)		
occurrences causally related to treatment / all	0 / 9		
deaths causally related to treatment / all	0 / 2		
Haemoptysis			
subjects affected / exposed	2 / 692 (0.29%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
Hydropneumothorax			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Hypoxia			
subjects affected / exposed	9 / 692 (1.30%)		
occurrences causally related to treatment / all	0 / 9		
deaths causally related to treatment / all	0 / 4		
Pleural effusion			
subjects affected / exposed	2 / 692 (0.29%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	1 / 2		
Pneumothorax			
subjects affected / exposed	3 / 692 (0.43%)		
occurrences causally related to treatment / all	1 / 3		
deaths causally related to treatment / all	0 / 0		
Pulmonary haemorrhage			
subjects affected / exposed	6 / 692 (0.87%)		
occurrences causally related to treatment / all	0 / 6		
deaths causally related to treatment / all	0 / 4		
Pulmonary oedema			
subjects affected / exposed	2 / 692 (0.29%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
Respiratory arrest			

subjects affected / exposed	2 / 692 (0.29%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 1		
Respiratory failure			
subjects affected / exposed	7 / 692 (1.01%)		
occurrences causally related to treatment / all	0 / 7		
deaths causally related to treatment / all	0 / 7		
Pulmonary alveolar haemorrhage			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Investigations			
Acoustic stimulation tests			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Blood creatinine			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Blood creatinine increased			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Cytomegalovirus test positive			
subjects affected / exposed	2 / 692 (0.29%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
Liver function test abnormal			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Oxygen saturation decreased			

subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Platelet count decreased			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Transaminases increased			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Injury, poisoning and procedural complications			
Collapse of lung			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Chemical peritonitis			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Cardiac disorders			
Arrhythmia			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Atrial fibrillation			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Bradycardia			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	1 / 1		

Cardiac arrest			
subjects affected / exposed	2 / 692 (0.29%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	1 / 2		
Cardiac failure			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	1 / 1		
Pericardial effusion			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	1 / 1		
Pericarditis			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Left ventricular dysfunction			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Nervous system disorders			
Cerebral haemorrhage			
subjects affected / exposed	2 / 692 (0.29%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	1 / 2		
Cerebral infarction			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 1		
Convulsion			
subjects affected / exposed	8 / 692 (1.16%)		
occurrences causally related to treatment / all	2 / 8		
deaths causally related to treatment / all	2 / 2		
Haemorrhage intracranial			



subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
Intracranial pressure increased			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Paraplegia			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Mental impairment			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Blood and lymphatic system disorders			
Anaemia			
subjects affected / exposed	3 / 692 (0.43%)		
occurrences causally related to treatment / all	1 / 3		
deaths causally related to treatment / all	0 / 1		
Disseminated intravascular coagulation			
subjects affected / exposed	5 / 692 (0.72%)		
occurrences causally related to treatment / all	1 / 5		
deaths causally related to treatment / all	1 / 2		
Febrile neutropenia			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Thrombocytopenia			
subjects affected / exposed	3 / 692 (0.43%)		
occurrences causally related to treatment / all	1 / 3		
deaths causally related to treatment / all	1 / 3		
Thrombotic microangiopathy			

subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Thrombotic thrombocytopenic purpura			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	1 / 1		
Gastrointestinal disorders			
Gastrointestinal haemorrhage			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Haematochezia			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Neutropenic colitis			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Abdominal pain upper			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Hepatobiliary disorders			
Cholecystitis acute			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Hepatic failure			
subjects affected / exposed	3 / 692 (0.43%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 2		

Hepatic function abnormal subjects affected / exposed	4 / 692 (0.58%)		
occurrences causally related to treatment / all	0 / 4		
deaths causally related to treatment / all	0 / 0		
Hepatorenal failure subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
Hepatosplenomegaly subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Hepatotoxicity subjects affected / exposed	2 / 692 (0.29%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	1 / 1		
Hyperbilirubinaemia subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Liver disorder subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
Skin and subcutaneous tissue disorders Rash subjects affected / exposed	2 / 692 (0.29%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
Renal and urinary disorders Azotaemia subjects affected / exposed	3 / 692 (0.43%)		
occurrences causally related to treatment / all	1 / 3		
deaths causally related to treatment / all	1 / 2		

Cystitis haemorrhagic			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Nephropathy toxic			
subjects affected / exposed	2 / 692 (0.29%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 1		
Renal failure			
subjects affected / exposed	3 / 692 (0.43%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 1		
Renal failure acute			
subjects affected / exposed	9 / 692 (1.30%)		
occurrences causally related to treatment / all	2 / 9		
deaths causally related to treatment / all	2 / 4		
Renal failure chronic			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	1 / 1		
Renal impairment			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Endocrine disorders			
Euthyroid sick syndrome			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Infections and infestations			
Anal abscess			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		

Aspergillosis				
subjects affected / exposed	6 / 692 (0.87%)			
occurrences causally related to treatment / all	0 / 6			
deaths causally related to treatment / all	0 / 5			
Brain abscess				
subjects affected / exposed	1 / 692 (0.14%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 1			
Bronchopulmonary aspergillosis				
subjects affected / exposed	1 / 692 (0.14%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Cellulitis				
subjects affected / exposed	1 / 692 (0.14%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 1			
Cytomegalovirus infection				
subjects affected / exposed	2 / 692 (0.29%)			
occurrences causally related to treatment / all	0 / 2			
deaths causally related to treatment / all	0 / 0			
Enterococcal sepsis				
subjects affected / exposed	2 / 692 (0.29%)			
occurrences causally related to treatment / all	0 / 2			
deaths causally related to treatment / all	0 / 0			
Fungal infection				
subjects affected / exposed	2 / 692 (0.29%)			
occurrences causally related to treatment / all	1 / 2			
deaths causally related to treatment / all	1 / 2			
Liver abscess				
subjects affected / exposed	1 / 692 (0.14%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Meningitis				

subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Oral herpes			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Pneumonia			
subjects affected / exposed	57 / 692 (8.24%)		
occurrences causally related to treatment / all	2 / 57		
deaths causally related to treatment / all	2 / 53		
Pneumonia fungal			
subjects affected / exposed	2 / 692 (0.29%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 2		
Pneumonia primary atypical			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
Sepsis			
subjects affected / exposed	29 / 692 (4.19%)		
occurrences causally related to treatment / all	2 / 29		
deaths causally related to treatment / all	2 / 25		
Septic shock			
subjects affected / exposed	44 / 692 (6.36%)		
occurrences causally related to treatment / all	2 / 44		
deaths causally related to treatment / all	0 / 35		
Enterococcal infection			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	1 / 1		
Metabolism and nutrition disorders			
Acidosis			

subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	1 / 1		
Hypernatraemia			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Metabolic acidosis			
subjects affected / exposed	3 / 692 (0.43%)		
occurrences causally related to treatment / all	1 / 3		
deaths causally related to treatment / all	1 / 2		
Hyperkalaemia			
subjects affected / exposed	1 / 692 (0.14%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		

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

<b>Non-serious adverse events</b>	Voriconazole		
Total subjects affected by non-serious adverse events			
subjects affected / exposed	94 / 692 (13.58%)		
Investigations			
Blood creatinine increased			
subjects affected / exposed	10 / 692 (1.45%)		
occurrences (all)	10		
Liver function test abnormal			
subjects affected / exposed	14 / 692 (2.02%)		
occurrences (all)	14		
Transaminases increased			
subjects affected / exposed	10 / 692 (1.45%)		
occurrences (all)	10		
Vascular disorders			
Hypertension			
subjects affected / exposed	7 / 692 (1.01%)		
occurrences (all)	7		

General disorders and administration site conditions Pyrexia subjects affected / exposed occurrences (all)	12 / 692 (1.73%) 12		
Gastrointestinal disorders Diarrhoea subjects affected / exposed occurrences (all)  Vomiting subjects affected / exposed occurrences (all)	9 / 692 (1.30%) 9  7 / 692 (1.01%) 7		
Hepatobiliary disorders Hyperbilirubinaemia subjects affected / exposed occurrences (all)	14 / 692 (2.02%) 14		
Respiratory, thoracic and mediastinal disorders Dyspnoea subjects affected / exposed occurrences (all)	8 / 692 (1.16%) 8		
Skin and subcutaneous tissue disorders Rash subjects affected / exposed occurrences (all)	7 / 692 (1.01%) 7		
Metabolism and nutrition disorders Hypokalaemia subjects affected / exposed occurrences (all)	10 / 692 (1.45%) 10		



## 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