



Clinical trial results:

A 24 month, randomized, controlled, study to evaluate the efficacy and safety of concentration-controlled everolimus plus reduced tacrolimus compared to standard tacrolimus in recipients of living donor liver transplants

Summary

EudraCT number	2010-024527-25
Trial protocol	IT DE
Global end of trial date	10 October 2017

Results information

Result version number	v1
This version publication date	21 October 2018
First version publication date	21 October 2018

Trial information

Trial identification

Sponsor protocol code	CRAD001H2307
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Additional study identifiers

ISRCTN number	-
ClinicalTrials.gov id (NCT number)	NCT01888432
WHO universal trial number (UTN)	-

Notes:

Sponsors

Sponsor organisation name	Novartis Pharma AG
Sponsor organisation address	CH-4002, Basel, Switzerland,
Public contact	Clinical Disclosure Office, Novartis Pharma AG, 41 613241111, novartis.email@novartis.com
Scientific contact	Clinical Disclosure Office, Novartis Pharma AG, 41 613241111, novartis.email@novartis.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	10 October 2017
Is this the analysis of the primary completion data?	No
Global end of trial reached?	Yes
Global end of trial date	10 October 2017
Was the trial ended prematurely?	No

Notes:

General information about the trial

Main objective of the trial:

To demonstrate comparable efficacy as measured by the composite efficacy failure of tBPAR, GL or death (D) with everolimus in combination with reduced tacrolimus compared to standard exposure tacrolimus, at 12 months post-transplantation, in living donor liver transplant recipients.

Protection of trial subjects:

The study was in compliance with the ethical principles derived from the Declaration of Helsinki and the International Conference on Harmonization (ICH) Good Clinical Practice (GCP) guidelines. All the local regulatory requirements pertinent to safety of trial subjects were also followed during the conduct of the trial.

Background therapy: -

Evidence for comparator: -

Actual start date of recruitment	25 September 2013
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	Taiwan: 79
Country: Number of subjects enrolled	Turkey: 14
Country: Number of subjects enrolled	United States: 23
Country: Number of subjects enrolled	Canada: 10
Country: Number of subjects enrolled	Egypt: 5
Country: Number of subjects enrolled	Germany: 5
Country: Number of subjects enrolled	India: 29
Country: Number of subjects enrolled	Italy: 2
Country: Number of subjects enrolled	Japan: 28
Country: Number of subjects enrolled	Korea, Republic of: 79
Country: Number of subjects enrolled	Russian Federation: 2
Country: Number of subjects enrolled	Saudi Arabia: 2
Country: Number of subjects enrolled	Singapore: 6
Worldwide total number of subjects	284
EEA total number of subjects	7

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)	284
From 65 to 84 years	0
85 years and over	0

Subject disposition

Recruitment

Recruitment details:

In all, 284 patients were randomized after transplantation to EVR+Reduced TAC group and TAC Control group. Two patients were not eligible and randomized in IRT by mistake and to whom no study medication was given, and so did not have their data included

Pre-assignment

Screening details:

A total of 494 patients were screened. Of these, 448 patients received a liver transplant and entered the run-in period.

Period 1

Period 1 title	12-month analysis (FAS)
Is this the baseline period?	No
Allocation method	Randomised - controlled
Blinding used	Not blinded

Arms

Are arms mutually exclusive?	No
Arm title	EVR+Reduced TAC

Arm description:

Everolimus + reduced tacrolimus ± corticosteroids

Arm type	Experimental
Investigational medicinal product name	Everolimus
Investigational medicinal product code	RAD001
Other name	
Pharmaceutical forms	Tablet
Routes of administration	Oral use, Intravenous use

Dosage and administration details:

as 1.0 mg, 0.75 and 0.50 mg tablets

Arm title	TAC Control
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Arm description:

Standard tacrolimus ± corticosteroids

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

Dosage and administration details:

0.5 mg, 1.0 mg and 5.0 mg capsules

Number of subjects in period 1	EVR+Reduced TAC	TAC Control
Started	142	142
Completed	131	133
Not completed	11	9
Adverse event, serious fatal	4	3
Physician decision	1	1
Consent withdrawn by subject	6	3
Lost to follow-up	-	2

Period 2

Period 2 title	24-month analysis (FAS)
Is this the baseline period?	Yes ^[1]
Allocation method	Randomised - controlled
Blinding used	Not blinded

Arms

Are arms mutually exclusive?	Yes
Arm title	EVR+Reduced TAC

Arm description:

Everolimus + reduced tacrolimus ± corticosteroids

Arm type	Experimental
Investigational medicinal product name	Everolimus
Investigational medicinal product code	RAD001
Other name	
Pharmaceutical forms	Tablet
Routes of administration	Oral use, Intravenous use

Dosage and administration details:

as 1.0 mg, 0.75 and 0.50 mg tablets

Arm title	TAC Control
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Arm description:

Standard tacrolimus ± corticosteroids

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

Dosage and administration details:

0.5 mg, 1.0 mg and 5.0 mg capsules

Notes:

[1] - Period 1 is not the baseline period. It is expected that period 1 will be the baseline period.

Justification: Period 1 is the 12-month mark and period 2 covers the complete trial period.

Number of subjects in period 2	EVR+Reduced TAC	TAC Control
Started	142	142
Completed	125	125
Not completed	17	17
Adverse event, serious fatal	8	4
Physician decision	2	4
Consent withdrawn by subject	7	6
Graft loss	-	1
Lost to follow-up	-	2

Baseline characteristics

Reporting groups

Reporting group title	EVR+Reduced TAC
Reporting group description: Everolimus + reduced tacrolimus ± corticosteroids	
Reporting group title	TAC Control
Reporting group description: Standard tacrolimus ± corticosteroids	

Reporting group values	EVR+Reduced TAC	TAC Control	Total
Number of subjects	142	142	284
Age categorical			
Units: Subjects			
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)	142	142	284
From 65-84 years	0	0	0
85 years and over	0	0	0
Age Continuous			
Units: Years			
arithmetic mean	54.2	52.7	-
standard deviation	± 8.95	± 10.41	-
Sex: Female, Male			
Units: Subjects			
Female	38	43	81
Male	104	99	203
Race/Ethnicity, Customized			
Units: Subjects			
Caucasian	30	30	60
Asian	111	112	223
Other	1	0	1

End points

End points reporting groups

Reporting group title	EVR+Reduced TAC
Reporting group description: Everolimus + reduced tacrolimus ± corticosteroids	
Reporting group title	TAC Control
Reporting group description: Standard tacrolimus ± corticosteroids	
Reporting group title	EVR+Reduced TAC
Reporting group description: Everolimus + reduced tacrolimus ± corticosteroids	
Reporting group title	TAC Control
Reporting group description: Standard tacrolimus ± corticosteroids	

Primary: Number of participants with composite efficacy failure of treated biopsy proven acute rejection, graft loss or death in everolimus with reduced tacrolimus group compared to standard tacrolimus

End point title	Number of participants with composite efficacy failure of treated biopsy proven acute rejection, graft loss or death in everolimus with reduced tacrolimus group compared to standard tacrolimus
End point description: Rate of composite efficacy failure of treated biopsy proven acute rejection (tBPAR ≥ RAI score 3), graft loss (GL) or death (D) in everolimus with reduced tacrolimus group compared to standard tacrolimus at 12 months	
End point type	Primary
End point timeframe: 12 months post transplantation	

End point values	EVR+Reduced TAC	TAC Control		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	142	142		
Units: Participants	7	8		

Statistical analyses

Statistical analysis title	12-month analysis (FAS)
Statistical analysis description: non-inferior efficacy failure of the reduced tacrolimus regimen to control by rejecting the null hypothesis.	
Comparison groups	EVR+Reduced TAC v TAC Control

Number of subjects included in analysis	284
Analysis specification	Pre-specified
Analysis type	
P-value	< 0.001 ^[1]
Method	Z-test
Parameter estimate	Kaplan-Meier
Point estimate	-0.7
Confidence interval	
level	90 %
sides	2-sided
lower limit	-5.2
upper limit	3.7

Notes:

[1] - Z-test p-value for non-inferiority test (non-inferiority margin = 12%) is for one-sided test and should be compared to 0.05 significance level.

Secondary: Renal function by estimated glomerular filtration rate (eGFR) from randomization

End point title	Renal function by estimated glomerular filtration rate (eGFR) from randomization
End point description:	Renal function (change in estimated glomerular filtration rate (eGFR)) from randomization to Month 12 post transplantation with everolimus (EVR) in combination with reduced tacrolimus (rTAC) compared to standard exposure tacrolimus (TAC) in living donor liver transplant recipients.
End point type	Secondary
End point timeframe:	Month 12

End point values	EVR+Reduced TAC	TAC Control		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	142	142		
Units: mL/min/1.73 m ²				
least squares mean (standard error)	-7.94 (± 1.839)	-12.09 (± 1.824)		

Statistical analyses

Statistical analysis title	12-month
Comparison groups	EVR+Reduced TAC v TAC Control
Number of subjects included in analysis	284
Analysis specification	Pre-specified
Analysis type	
P-value	< 0.001
Method	ANCOVA
Parameter estimate	Mean difference (net)
Point estimate	4.15

Confidence interval	
level	90 %
sides	2-sided
lower limit	-0.09
upper limit	8.4
Variability estimate	Standard error of the mean
Dispersion value	2.574

Secondary: Compare renal function over time assessed by the change by eGFR, post-randomization

End point title	Compare renal function over time assessed by the change by eGFR, post-randomization
End point description: Compare evolution of post-randomization renal function over time assessed by the change in estimated GFR (MDRD-4), including changes from randomization to Months 12 and 24. Rate of change of renal function.	
End point type	Secondary
End point timeframe: Month 24	

End point values	EVR+Reduced TAC	TAC Control		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	142	142		
Units: Participants				
least squares mean (standard error)	-11.01 (\pm 1.928)	-14.26 (\pm 1.914)		

Statistical analyses

Statistical analysis title	12-month
Comparison groups	EVR+Reduced TAC v TAC Control
Number of subjects included in analysis	284
Analysis specification	Pre-specified
Analysis type	
P-value	< 0.001
Method	ANCOVA
Parameter estimate	Mean difference (final values)
Point estimate	3.25
Confidence interval	
level	90 %
sides	2-sided
lower limit	-1.21
upper limit	7.7
Variability estimate	Standard error of the mean
Dispersion value	2.699

Secondary: Compare incidence of a composite of tBPAR, graft loss, and death

End point title	Compare incidence of a composite of tBPAR, graft loss, and death
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End point description:

Compare between the treatment group EVR with rTAC vs standard TAC: incidence of a composite of tBPAR, graft loss, death

End point type	Secondary
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End point timeframe:

Month 24 post transplantation

End point values	EVR+Reduced TAC	TAC Control		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	142	142		
Units: Participant				
tBPAR/graft loss/death	12	11		
On-treatment tBPAR/graft loss/death	7	9		

Statistical analyses

No statistical analyses for this end point

Secondary: Compare incidence of tBPAR

End point title	Compare incidence of tBPAR
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End point description:

Compare between the treatment group EVR with rTAC vs standard TAC: Incidence of tBPAR

End point type	Secondary
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End point timeframe:

Month 12 and Month 24 post transplantation

End point values	EVR+Reduced TAC	TAC Control		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	142	142		
Units: Participants				
tBPAR - month 12	3	5		
tBPAR - month 24	4	6		
On-treatment tBPAR - month 24	3	6		

Statistical analyses

Statistical analysis title	Month 12
Comparison groups	EVR+Reduced TAC v TAC Control
Number of subjects included in analysis	284
Analysis specification	Pre-specified
Analysis type	
Parameter estimate	Kaplan-Meier
Point estimate	-1.4
Confidence interval	
level	90 %
sides	2-sided
lower limit	-4.7
upper limit	2

Statistical analysis title	Month 24 (tBPAR)
Comparison groups	EVR+Reduced TAC v TAC Control
Number of subjects included in analysis	284
Analysis specification	Pre-specified
Analysis type	
Parameter estimate	Kaplan-Meier
Point estimate	-1.2
Confidence interval	
level	90 %
sides	2-sided
lower limit	-5.1
upper limit	2.6

Statistical analysis title	Month 24 (on-treatment tBPAR)
Comparison groups	EVR+Reduced TAC v TAC Control
Number of subjects included in analysis	284
Analysis specification	Pre-specified
Analysis type	
Parameter estimate	Kaplan-Meier
Point estimate	-2.1
Confidence interval	
level	90 %
sides	2-sided
lower limit	-5.7
upper limit	1.4

Secondary: Compare incidence of BPAR

End point title	Compare incidence of BPAR
End point description:	Compare between the treatment group EVR with rTAC vs standard TAC: incidence of a composite of biopsy proven acute rejection (BPAR)
End point type	Secondary
End point timeframe:	Month 12 and Month 24 post transplantation

End point values	EVR+Reduced TAC	TAC Control		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	142	142		
Units: Participants				
Month 12	7	6		
Month 24	8	7		

Statistical analyses

Statistical analysis title	Month 12
Comparison groups	EVR+Reduced TAC v TAC Control
Number of subjects included in analysis	284
Analysis specification	Pre-specified
Analysis type	
Parameter estimate	Kaplan-Meier
Point estimate	0.9
Confidence interval	
level	90 %
sides	2-sided
lower limit	-3.4
upper limit	5.1

Statistical analysis title	Month 24
Comparison groups	EVR+Reduced TAC v TAC Control
Number of subjects included in analysis	284
Analysis specification	Pre-specified
Analysis type	
Parameter estimate	Kaplan-Meier
Point estimate	1

Confidence interval	
level	90 %
sides	2-sided
lower limit	-3.6
upper limit	5.6

Secondary: Compare incidence of graft loss

End point title	Compare incidence of graft loss
End point description:	Compare between the treatment group EVR with rTAC vs standard TAC: incidence of graft loss
End point type	Secondary
End point timeframe:	Month 12 and Month 24 post transplantation

End point values	EVR+Reduced TAC	TAC Control		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	142	142		
Units: Participants				
Month 12	0	0		
month 24	0	1		
month 24 (on-treatment graft loss)	0	0		

Statistical analyses

Statistical analysis title	graft loss at month 24
Comparison groups	EVR+Reduced TAC v TAC Control
Number of subjects included in analysis	284
Analysis specification	Pre-specified
Analysis type	
Parameter estimate	Kaplan-Meier
Point estimate	-0.8
Confidence interval	
level	90 %
sides	2-sided
lower limit	-2.1
upper limit	0.5

Secondary: Compare incidence of a composite of death or graft loss

End point title	Compare incidence of a composite of death or graft loss
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End point description:

Compare between the treatment group EVR with rTAC vs standard TAC: Incidence of a composite of death or graft loss

End point type Secondary

End point timeframe:

Month 12 and Month 24 post transplantation

End point values	EVR+Reduced TAC	TAC Control		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	142	142		
Units: Participants				
Month 12	4	3		
Month 24	8	5		

Statistical analyses

Statistical analysis title	Month 24
Comparison groups	EVR+Reduced TAC v TAC Control
Number of subjects included in analysis	284
Analysis specification	Pre-specified
Analysis type	
Parameter estimate	Kaplan-Meier
Point estimate	2.3
Confidence interval	
level	90 %
sides	2-sided
lower limit	-2.1
upper limit	6.6

Statistical analysis title	Month 12
Comparison groups	EVR+Reduced TAC v TAC Control
Number of subjects included in analysis	284
Analysis specification	Pre-specified
Analysis type	
Parameter estimate	Kaplan-Meier
Point estimate	0.7
Confidence interval	
level	90 %
sides	2-sided
lower limit	-2.5
upper limit	3.8

Secondary: Compare incidence of death

End point title	Compare incidence of death
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End point description:

Compare between the treatment group EVR with rTAC vs standard TAC: incidence of death

End point type	Secondary
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End point timeframe:

Month 12 and Month 24 post transplantation

End point values	EVR+Reduced TAC	TAC Control		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	142	142		
Units: Participants				
Month 12	4	3		
Month 24	8	4		
Month 24 (on-treatment death)	4	3		

Statistical analyses

Statistical analysis title	Month 12
Comparison groups	EVR+Reduced TAC v TAC Control
Number of subjects included in analysis	284
Analysis specification	Pre-specified
Analysis type	
Parameter estimate	Kaplan-Meier
Point estimate	0.7
Confidence interval	
level	90 %
sides	2-sided
lower limit	-2.5
upper limit	3.8

Statistical analysis title	Month 24
Comparison groups	EVR+Reduced TAC v TAC Control

Number of subjects included in analysis	284
Analysis specification	Pre-specified
Analysis type	
Parameter estimate	Kaplan-Meier
Point estimate	3
Confidence interval	
level	90 %
sides	2-sided
lower limit	-1.1
upper limit	7.2

Statistical analysis title	Month 24 (On-treatment death)
Comparison groups	EVR+Reduced TAC v TAC Control
Number of subjects included in analysis	284
Analysis specification	Pre-specified
Analysis type	
Parameter estimate	Kaplan-Meier
Point estimate	0.7
Confidence interval	
level	90 %
sides	2-sided
lower limit	-2.5
upper limit	3.9

Secondary: Compare incidence of AR

End point title	Compare incidence of AR
End point description: Compare between the treatment group EVR with rTAC vs standard TAC: incidence of acute rejection (AR)	
End point type	Secondary
End point timeframe: Month 12 and Month 24 post transplantation	

End point values	EVR+Reduced TAC	TAC Control		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	142	142		
Units: Participants				
Month 12	9	8		
Month 24	12	9		

Statistical analyses

Statistical analysis title	Incidence rate of AR - month 12
Comparison groups	EVR+Reduced TAC v TAC Control
Number of subjects included in analysis	284
Analysis specification	Pre-specified
Analysis type	
Parameter estimate	Risk difference (RD)
Point estimate	0.7
Confidence interval	
level	90 %
sides	2-sided
lower limit	-3.9
upper limit	5.3

Statistical analysis title	Incidence rate of AR - month 24
Comparison groups	EVR+Reduced TAC v TAC Control
Number of subjects included in analysis	284
Analysis specification	Pre-specified
Analysis type	
Parameter estimate	Risk difference (RD)
Point estimate	2.1
Confidence interval	
level	90 %
sides	2-sided
lower limit	-3
upper limit	7.2

Secondary: Compare incidence of tAR

End point title	Compare incidence of tAR
End point description:	Compare between the treatment group EVR with rTAC vs standard TAC: incidence of treated acute rejection (tAR).
End point type	Secondary
End point timeframe:	Month 12 and Month 24 post transplantation

End point values	EVR+Reduced TAC	TAC Control		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	142	142		
Units: Participants				
Month 12	5	6		
Month 24	7	7		

Statistical analyses

Statistical analysis title	Incidence rate of tAR - month 12
Comparison groups	EVR+Reduced TAC v TAC Control
Number of subjects included in analysis	284
Analysis specification	Pre-specified
Analysis type	
Parameter estimate	Risk ratio (RR)
Point estimate	-0.7
Confidence interval	
level	90 %
sides	2-sided
lower limit	-4.5
upper limit	3.1

Statistical analysis title	Incidence rate of tAR - month 24
Comparison groups	EVR+Reduced TAC v TAC Control
Number of subjects included in analysis	284
Analysis specification	Pre-specified
Analysis type	
Parameter estimate	Risk ratio (RR)
Point estimate	0
Confidence interval	
level	90 %
sides	2-sided
lower limit	-4.2
upper limit	4.2

Secondary: Compare time to recurrence of HCC in subjects with a diagnosis of HCC at the time of liver transplantation

End point title	Compare time to recurrence of HCC in subjects with a diagnosis of HCC at the time of liver transplantation
End point description:	Patients transplanted for HCC or with HCC diagnosed at time of transplantation were monitored for HCC recurrence according to local practice. For example routine laboratory monitoring/tests, tumor markers, hepatic ultrasound, computed tomography scans (CAT, CT) or MRI (especially Fe-MRI) on a regular basis per local practice.
End point type	Secondary
End point timeframe:	Month 12 and Month 24

End point values	EVR+Reduced TAC	TAC Control		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	142	141		
Units: Participants				
HCC recurrence (n/M) - month 12	0	5		
HCC recurrence (n/M) - month 24	1	6		

Statistical analyses

No statistical analyses for this end point

Secondary: Number of subjects experiencing adverse events/infections by SOC

End point title	Number of subjects experiencing adverse events/infections by SOC
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End point description:

End point type	Secondary
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End point timeframe:

Month 24

End point values	EVR+Reduced TAC	TAC Control		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	142	141		
Units: Participants				
Any AE/infection	140	136		
Blood and lymphatic system disorders	44	32		
Cardiac disorders	15	12		
Congenital, familial and genetic disorders	1	2		
Ear and labyrinth disorders	2	5		
Endocrine disorders	1	2		
Eye disorders	17	15		
Gastrointestinal disorders	98	74		
General disorders&admin site conditions	48	42		
Hepatobiliary disorders	44	40		
Immune system disorders	8	11		
Infections and infestations	84	70		
Injury, poisoning&proced. complications	36	28		
Investigations	61	68		
Metabolism and nutrition disorders	87	60		

Musculoskeletal and connective tissue disorders	30	43		
Neo benign, malig&unspecified (cysts&polyps)	10	17		
Nervous system disorders	38	44		
Product issues#	1	1		
Psychiatric disorders	33	26		
Renal and urinary disorders	46	36		
Reproductive system&breast dis.	9	12		
Respiratory, thoracic&mediastinal dis.	34	39		
Skin&subcutaneous tissue disorders	39	44		
Vascular disorders	38	30		

Statistical analyses

No statistical analyses for this end point

Secondary: Compare incidence of notable safety events (SAEs, infections and serious infections leading to premature discontinuation)

End point title	Compare incidence of notable safety events (SAEs, infections and serious infections leading to premature discontinuation)
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End point description:

Notable events include death, Serious AE/infection,, and AE/infection leading to discontinuation of study medication.

End point type	Secondary
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End point timeframe:

Month 24

End point values	EVR+Reduced TAC	TAC Control		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	142	141		
Units: Participants				
Any notable events	86	82		
Death	8	4		
Serious AE/Infection	83	78		
AE/Infection lead. to premature disc of study med	21	18		

Statistical analyses

No statistical analyses for this end point

Adverse events

Adverse events information

Timeframe for reporting adverse events:

Adverse Events (AEs) are collected from First Patient First Visit (FPFV) until Last Patient Last Visit (LPLV). All AEs reported in this record are from date of First Patient First Treatment until Last Patient Last Visit) up to approximately 4 years.

Assessment type	Systematic
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Dictionary used

Dictionary name	MedDRA
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Dictionary version	20.1
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Reporting groups

Reporting group title	EVR+Reduced TAC
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Reporting group description:

EVR+Reduced TAC

Reporting group title	TAC Control
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Reporting group description:

TAC Control

Serious adverse events	EVR+Reduced TAC	TAC Control	
Total subjects affected by serious adverse events			
subjects affected / exposed	83 / 142 (58.45%)	78 / 141 (55.32%)	
number of deaths (all causes)	8	4	
number of deaths resulting from adverse events	0	0	
Neoplasms benign, malignant and unspecified (incl cysts and polyps)			
Adenocarcinoma of colon			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Bone giant cell tumour			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hepatic angiosarcoma			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hepatic neoplasm			

subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Hepatocellular carcinoma		
subjects affected / exposed	1 / 142 (0.70%)	4 / 141 (2.84%)
occurrences causally related to treatment / all	0 / 1	0 / 6
deaths causally related to treatment / all	0 / 1	0 / 1
Lipofibroma		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Lung neoplasm malignant		
subjects affected / exposed	2 / 142 (1.41%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 2	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Metastases to bone		
subjects affected / exposed	0 / 142 (0.00%)	2 / 141 (1.42%)
occurrences causally related to treatment / all	0 / 0	0 / 3
deaths causally related to treatment / all	0 / 0	0 / 1
Metastases to central nervous system		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Metastases to lung		
subjects affected / exposed	0 / 142 (0.00%)	5 / 141 (3.55%)
occurrences causally related to treatment / all	0 / 0	1 / 6
deaths causally related to treatment / all	0 / 0	0 / 1
Metastases to spine		
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Prostate cancer		

subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Squamous cell carcinoma of the hypopharynx			
subjects affected / exposed	1 / 142 (0.70%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 1	0 / 1	
deaths causally related to treatment / all	0 / 1	0 / 0	
Squamous cell carcinoma of the oral cavity			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 1	
Vascular disorders			
Arterial haemorrhage			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Haematoma			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hot flush			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hypotension			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Lymphangiopathy			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Vena cava thrombosis			

subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Venous stenosis			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
General disorders and administration site conditions			
Chest discomfort			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Chills			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Crepitations			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hyperpyrexia			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Incarcerated hernia			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Multiple organ dysfunction syndrome			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 1	0 / 0	
Non-cardiac chest pain			

subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pyrexia			
subjects affected / exposed	11 / 142 (7.75%)	8 / 141 (5.67%)	
occurrences causally related to treatment / all	2 / 15	0 / 8	
deaths causally related to treatment / all	0 / 0	0 / 0	
Immune system disorders			
Liver transplant rejection			
subjects affected / exposed	3 / 142 (2.11%)	3 / 141 (2.13%)	
occurrences causally related to treatment / all	1 / 3	0 / 3	
deaths causally related to treatment / all	0 / 0	0 / 0	
Transplant rejection			
subjects affected / exposed	2 / 142 (1.41%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 2	1 / 2	
deaths causally related to treatment / all	0 / 0	0 / 0	
Respiratory, thoracic and mediastinal disorders			
Acute pulmonary oedema			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Acute respiratory failure			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Cough			
subjects affected / exposed	1 / 142 (0.70%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 1	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Dyspnoea			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	

Hypoxia			
subjects affected / exposed	1 / 142 (0.70%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 1	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pharyngeal ulceration			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pleural effusion			
subjects affected / exposed	1 / 142 (0.70%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 1	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pneumonia aspiration			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pneumothorax			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pulmonary embolism			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pulmonary infarction			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 1	
Pulmonary mass			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Respiratory failure			

subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 1	0 / 0	
Wheezing			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Psychiatric disorders			
Alcoholism			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Insomnia			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	1 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Mental status changes			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Suicidal behaviour			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Investigations			
Alanine aminotransferase increased			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Aspartate aminotransferase increased			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	

Blood alkaline phosphatase increased			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
C-reactive protein increased			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Gamma-glutamyltransferase increased			
subjects affected / exposed	0 / 142 (0.00%)	2 / 141 (1.42%)	
occurrences causally related to treatment / all	0 / 0	1 / 2	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hepatic enzyme increased			
subjects affected / exposed	4 / 142 (2.82%)	2 / 141 (1.42%)	
occurrences causally related to treatment / all	0 / 7	1 / 4	
deaths causally related to treatment / all	0 / 0	0 / 0	
Immunosuppressant drug level increased			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	1 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Liver function test increased			
subjects affected / exposed	1 / 142 (0.70%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	1 / 1	1 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Transaminases increased			
subjects affected / exposed	1 / 142 (0.70%)	2 / 141 (1.42%)	
occurrences causally related to treatment / all	0 / 1	1 / 2	
deaths causally related to treatment / all	0 / 0	0 / 0	
Injury, poisoning and procedural complications			
Anastomotic stenosis			

subjects affected / exposed	2 / 142 (1.41%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	1 / 3	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Biliary anastomosis complication		
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 0	0 / 2
deaths causally related to treatment / all	0 / 0	0 / 0
Burns second degree		
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Chemical peritonitis		
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Contusion		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Facial bones fracture		
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Femoral neck fracture		
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Graft loss		
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Incarcerated incisional hernia		

subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Incision site pain		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Incisional hernia		
subjects affected / exposed	5 / 142 (3.52%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 5	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Multiple injuries		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Post procedural bile leak		
subjects affected / exposed	1 / 142 (0.70%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	1 / 1	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Post procedural haemorrhage		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Post procedural inflammation		
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Procedural complication		
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Procedural pain		

subjects affected / exposed	1 / 142 (0.70%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 1	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Spinal compression fracture			
subjects affected / exposed	2 / 142 (1.41%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 2	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Spinal fracture			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Wound dehiscence			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Cardiac disorders			
Acute myocardial infarction			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Angina pectoris			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Atrial fibrillation			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Bradycardia			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 1	0 / 0	
Cardiac arrest			

subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 1	0 / 0	
Cardiac discomfort			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Cardiopulmonary failure			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 1	0 / 0	
Coronary artery disease			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Mitral valve incompetence			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Supraventricular tachycardia			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Nervous system disorders			
Altered state of consciousness			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 1	
Dizziness			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Headache			

subjects affected / exposed	0 / 142 (0.00%)	2 / 141 (1.42%)	
occurrences causally related to treatment / all	0 / 0	1 / 2	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hypoaesthesia			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 2	
deaths causally related to treatment / all	0 / 0	0 / 0	
Migraine			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Neuropathy peripheral			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Restless legs syndrome			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Seizure			
subjects affected / exposed	1 / 142 (0.70%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 1	1 / 2	
deaths causally related to treatment / all	0 / 1	0 / 0	
Tremor			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Wernicke's encephalopathy			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Blood and lymphatic system disorders			
Anaemia			

subjects affected / exposed	2 / 142 (1.41%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	2 / 3	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Leukocytosis			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Leukopenia			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Thrombocytopenia			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Thrombotic microangiopathy			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Eye disorders			
Cataract			
subjects affected / exposed	1 / 142 (0.70%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 1	0 / 2	
deaths causally related to treatment / all	0 / 0	0 / 0	
Lens dislocation			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Gastrointestinal disorders			
Abdominal hernia			
subjects affected / exposed	3 / 142 (2.11%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 3	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Abdominal incarcerated hernia			

subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Abdominal pain		
subjects affected / exposed	7 / 142 (4.93%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 7	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Abdominal pain upper		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Anal incontinence		
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Ascites		
subjects affected / exposed	0 / 142 (0.00%)	2 / 141 (1.42%)
occurrences causally related to treatment / all	0 / 0	0 / 3
deaths causally related to treatment / all	0 / 0	0 / 0
Colitis		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Constipation		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Crohn's disease		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Diarrhoea		

subjects affected / exposed	2 / 142 (1.41%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 2	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Duodenal perforation		
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Duodenal ulcer		
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Duodenal ulcer haemorrhage		
subjects affected / exposed	1 / 142 (0.70%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	1 / 1	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Enteritis		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Gastric ulcer haemorrhage		
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Gastrointestinal inflammation		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Haemorrhoids		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Ileus		

subjects affected / exposed	1 / 142 (0.70%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 1	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Ileus paralytic			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Incarcerated umbilical hernia			
subjects affected / exposed	1 / 142 (0.70%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 1	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Inguinal hernia			
subjects affected / exposed	0 / 142 (0.00%)	2 / 141 (1.42%)	
occurrences causally related to treatment / all	0 / 0	0 / 2	
deaths causally related to treatment / all	0 / 0	0 / 0	
Intra-abdominal fluid collection			
subjects affected / exposed	1 / 142 (0.70%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 1	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Leukoplakia oral			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Mesenteric haemorrhage			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 1	0 / 0	
Nausea			
subjects affected / exposed	1 / 142 (0.70%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 1	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pancreatitis			

subjects affected / exposed	1 / 142 (0.70%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 1	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pancreatitis acute			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 3	
deaths causally related to treatment / all	0 / 0	0 / 0	
Parotid gland enlargement			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Peptic ulcer perforation			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pouchitis			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 2	
deaths causally related to treatment / all	0 / 0	0 / 0	
Small intestinal haemorrhage			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 1	0 / 0	
Small intestine ulcer			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Strangulated umbilical hernia			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Vomiting			

subjects affected / exposed	2 / 142 (1.41%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 2	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hepatobiliary disorders			
Autoimmune hepatitis			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Bile duct obstruction			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Bile duct stenosis			
subjects affected / exposed	10 / 142 (7.04%)	6 / 141 (4.26%)	
occurrences causally related to treatment / all	0 / 19	0 / 9	
deaths causally related to treatment / all	0 / 0	0 / 0	
Bile duct stone			
subjects affected / exposed	2 / 142 (1.41%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 2	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Biliary cirrhosis primary			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Biliary dilatation			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 2	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Biliary fistula			
subjects affected / exposed	2 / 142 (1.41%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 2	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Biloma			

subjects affected / exposed	1 / 142 (0.70%)	2 / 141 (1.42%)
occurrences causally related to treatment / all	0 / 1	0 / 2
deaths causally related to treatment / all	0 / 0	0 / 0
Cholangitis		
subjects affected / exposed	8 / 142 (5.63%)	9 / 141 (6.38%)
occurrences causally related to treatment / all	2 / 12	0 / 14
deaths causally related to treatment / all	0 / 0	0 / 0
Cholangitis acute		
subjects affected / exposed	2 / 142 (1.41%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 4	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Cholelithiasis		
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Cholelithiasis obstructive		
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Dilatation intrahepatic duct acquired		
subjects affected / exposed	1 / 142 (0.70%)	2 / 141 (1.42%)
occurrences causally related to treatment / all	0 / 1	0 / 2
deaths causally related to treatment / all	0 / 0	0 / 0
Hepatic artery thrombosis		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	1 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Hepatic function abnormal		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	1 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Hepatic steatosis		

subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hepatitis cholestatic			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Jaundice			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Jaundice cholestatic			
subjects affected / exposed	0 / 142 (0.00%)	2 / 141 (1.42%)	
occurrences causally related to treatment / all	0 / 0	0 / 3	
deaths causally related to treatment / all	0 / 0	0 / 0	
Portal vein stenosis			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Portal vein thrombosis			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Skin and subcutaneous tissue disorders			
Skin ulcer			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Renal and urinary disorders			
Acute kidney injury			
subjects affected / exposed	2 / 142 (1.41%)	2 / 141 (1.42%)	
occurrences causally related to treatment / all	1 / 2	0 / 2	
deaths causally related to treatment / all	0 / 0	0 / 0	
Chromaturia			

subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Chronic kidney disease			
subjects affected / exposed	2 / 142 (1.41%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	2 / 2	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Haematuria			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Proteinuria			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Renal failure			
subjects affected / exposed	2 / 142 (1.41%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 2	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Renal impairment			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	1 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Musculoskeletal and connective tissue disorders			
Back pain			
subjects affected / exposed	2 / 142 (1.41%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 2	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Intervertebral disc protrusion			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Lumbar spinal stenosis			

subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Musculoskeletal pain			
subjects affected / exposed	0 / 142 (0.00%)	2 / 141 (1.42%)	
occurrences causally related to treatment / all	0 / 0	0 / 2	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pain in extremity			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Spondylolisthesis			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Infections and infestations			
Abdominal abscess			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Appendicitis			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Arteriovenous graft site infection			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Bacterial sepsis			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Biliary tract infection			

subjects affected / exposed	1 / 142 (0.70%)	2 / 141 (1.42%)
occurrences causally related to treatment / all	0 / 1	0 / 3
deaths causally related to treatment / all	0 / 0	0 / 0
Bronchopulmonary aspergillosis		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Cellulitis		
subjects affected / exposed	3 / 142 (2.11%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	1 / 3	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Clostridium difficile colitis		
subjects affected / exposed	3 / 142 (2.11%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	2 / 3	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Cytomegalovirus viraemia		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	1 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Diarrhoea infectious		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Enterococcal bacteraemia		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	1 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Epididymitis		
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Fungal sepsis		

subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 1	0 / 0
Gastroenteritis		
subjects affected / exposed	4 / 142 (2.82%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	1 / 4	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Gastroenteritis norovirus		
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Groin abscess		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	1 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Hepatitis C		
subjects affected / exposed	5 / 142 (3.52%)	3 / 141 (2.13%)
occurrences causally related to treatment / all	2 / 5	0 / 3
deaths causally related to treatment / all	0 / 0	0 / 1
Hepatitis E		
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Herpes zoster		
subjects affected / exposed	1 / 142 (0.70%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	1 / 1	1 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Infection		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Liver abscess		

subjects affected / exposed	2 / 142 (1.41%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 2	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Lower respiratory tract infection		
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 1
Orchitis		
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Peritonitis		
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Pharyngitis		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Pneumonia		
subjects affected / exposed	10 / 142 (7.04%)	5 / 141 (3.55%)
occurrences causally related to treatment / all	2 / 11	1 / 6
deaths causally related to treatment / all	0 / 2	0 / 0
Pneumonia legionella		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	1 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Pneumonia viral		
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 0	1 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Post procedural cellulitis		

subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Post procedural sepsis		
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 0	1 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Postoperative wound infection		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Prostatic abscess		
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
occurrences causally related to treatment / all	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0
Pulmonary tuberculosis		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Pyelonephritis acute		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	1 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Sepsis		
subjects affected / exposed	1 / 142 (0.70%)	2 / 141 (1.42%)
occurrences causally related to treatment / all	0 / 1	0 / 2
deaths causally related to treatment / all	0 / 0	0 / 0
Septic shock		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 1	0 / 0
Staphylococcal bacteraemia		

subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Systemic infection			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Tuberculosis			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Upper respiratory tract infection			
subjects affected / exposed	3 / 142 (2.11%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	1 / 3	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Upper respiratory tract infection bacterial			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	1 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Urinary tract infection			
subjects affected / exposed	1 / 142 (0.70%)	3 / 141 (2.13%)	
occurrences causally related to treatment / all	0 / 1	0 / 3	
deaths causally related to treatment / all	0 / 0	0 / 0	
Urosepsis			
subjects affected / exposed	2 / 142 (1.41%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	1 / 2	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Metabolism and nutrition disorders			
Diabetes mellitus			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Diabetes mellitus inadequate control			

subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Fluid overload			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Gout			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hyperglycaemia			
subjects affected / exposed	1 / 142 (0.70%)	3 / 141 (2.13%)	
occurrences causally related to treatment / all	0 / 1	1 / 3	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hyperkalaemia			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hypertriglyceridaemia			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hypoalbuminaemia			
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hypokalaemia			
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hyponatraemia			

subjects affected / exposed	2 / 142 (1.41%)	2 / 141 (1.42%)
occurrences causally related to treatment / all	0 / 2	0 / 4
deaths causally related to treatment / all	0 / 0	0 / 0
Malnutrition		
subjects affected / exposed	1 / 142 (0.70%)	0 / 141 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0
Obesity		
subjects affected / exposed	0 / 142 (0.00%)	1 / 141 (0.71%)
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: 5 %

Non-serious adverse events	EVR+Reduced TAC	TAC Control	
Total subjects affected by non-serious adverse events			
subjects affected / exposed	125 / 142 (88.03%)	120 / 141 (85.11%)	
Vascular disorders			
Hypertension			
subjects affected / exposed	29 / 142 (20.42%)	23 / 141 (16.31%)	
occurrences (all)	31	25	
General disorders and administration site conditions			
Oedema peripheral			
subjects affected / exposed	16 / 142 (11.27%)	8 / 141 (5.67%)	
occurrences (all)	17	11	
Pyrexia			
subjects affected / exposed	18 / 142 (12.68%)	20 / 141 (14.18%)	
occurrences (all)	40	29	
Respiratory, thoracic and mediastinal disorders			
Cough			
subjects affected / exposed	12 / 142 (8.45%)	15 / 141 (10.64%)	
occurrences (all)	14	16	
Psychiatric disorders			
Depression			

subjects affected / exposed occurrences (all)	2 / 142 (1.41%) 2	8 / 141 (5.67%) 8	
Insomnia subjects affected / exposed occurrences (all)	26 / 142 (18.31%) 31	13 / 141 (9.22%) 17	
Investigations Alanine aminotransferase increased subjects affected / exposed occurrences (all)	5 / 142 (3.52%) 5	8 / 141 (5.67%) 8	
Blood creatinine increased subjects affected / exposed occurrences (all)	10 / 142 (7.04%) 11	13 / 141 (9.22%) 21	
Hepatic enzyme abnormal subjects affected / exposed occurrences (all)	8 / 142 (5.63%) 13	19 / 141 (13.48%) 36	
Hepatic enzyme increased subjects affected / exposed occurrences (all)	16 / 142 (11.27%) 21	19 / 141 (13.48%) 29	
Nervous system disorders Dizziness subjects affected / exposed occurrences (all)	3 / 142 (2.11%) 3	9 / 141 (6.38%) 9	
Headache subjects affected / exposed occurrences (all)	19 / 142 (13.38%) 21	14 / 141 (9.93%) 18	
Tremor subjects affected / exposed occurrences (all)	4 / 142 (2.82%) 4	11 / 141 (7.80%) 11	
Blood and lymphatic system disorders Anaemia subjects affected / exposed occurrences (all)	18 / 142 (12.68%) 20	15 / 141 (10.64%) 20	
Leukopenia subjects affected / exposed occurrences (all)	16 / 142 (11.27%) 34	7 / 141 (4.96%) 8	
Thrombocytopenia			

subjects affected / exposed occurrences (all)	14 / 142 (9.86%) 14	3 / 141 (2.13%) 4	
Eye disorders Cataract subjects affected / exposed occurrences (all)	8 / 142 (5.63%) 8	4 / 141 (2.84%) 4	
Gastrointestinal disorders Abdominal pain subjects affected / exposed occurrences (all)	19 / 142 (13.38%) 19	12 / 141 (8.51%) 15	
Constipation subjects affected / exposed occurrences (all)	13 / 142 (9.15%) 18	16 / 141 (11.35%) 16	
Diarrhoea subjects affected / exposed occurrences (all)	32 / 142 (22.54%) 41	19 / 141 (13.48%) 28	
Nausea subjects affected / exposed occurrences (all)	7 / 142 (4.93%) 7	9 / 141 (6.38%) 9	
Mouth ulceration subjects affected / exposed occurrences (all)	17 / 142 (11.97%) 18	6 / 141 (4.26%) 7	
Stomatitis subjects affected / exposed occurrences (all)	19 / 142 (13.38%) 25	5 / 141 (3.55%) 6	
Vomiting subjects affected / exposed occurrences (all)	7 / 142 (4.93%) 12	10 / 141 (7.09%) 11	
Hepatobiliary disorders Hepatic steatosis subjects affected / exposed occurrences (all)	10 / 142 (7.04%) 10	7 / 141 (4.96%) 7	
Skin and subcutaneous tissue disorders Pruritus subjects affected / exposed occurrences (all)	22 / 142 (15.49%) 26	18 / 141 (12.77%) 22	
Rash			

subjects affected / exposed occurrences (all)	7 / 142 (4.93%) 11	10 / 141 (7.09%) 10	
Renal and urinary disorders			
Renal failure subjects affected / exposed occurrences (all)	3 / 142 (2.11%) 3	13 / 141 (9.22%) 15	
Renal impairment subjects affected / exposed occurrences (all)	11 / 142 (7.75%) 12	5 / 141 (3.55%) 7	
Musculoskeletal and connective tissue disorders			
Arthralgia subjects affected / exposed occurrences (all)	5 / 142 (3.52%) 5	9 / 141 (6.38%) 9	
Back pain subjects affected / exposed occurrences (all)	8 / 142 (5.63%) 8	10 / 141 (7.09%) 10	
Infections and infestations			
Nasopharyngitis subjects affected / exposed occurrences (all)	20 / 142 (14.08%) 38	16 / 141 (11.35%) 22	
Upper respiratory tract infection subjects affected / exposed occurrences (all)	18 / 142 (12.68%) 25	11 / 141 (7.80%) 17	
Metabolism and nutrition disorders			
Decreased appetite subjects affected / exposed occurrences (all)	9 / 142 (6.34%) 10	13 / 141 (9.22%) 14	
Dyslipidaemia subjects affected / exposed occurrences (all)	13 / 142 (9.15%) 14	4 / 141 (2.84%) 5	
Hypercholesterolaemia subjects affected / exposed occurrences (all)	22 / 142 (15.49%) 31	2 / 141 (1.42%) 2	
Hyperkalaemia subjects affected / exposed occurrences (all)	20 / 142 (14.08%) 25	13 / 141 (9.22%) 21	

Hyperlipidaemia		
subjects affected / exposed	22 / 142 (15.49%)	8 / 141 (5.67%)
occurrences (all)	23	8
Hyperuricaemia		
subjects affected / exposed	7 / 142 (4.93%)	8 / 141 (5.67%)
occurrences (all)	9	11
Hypokalaemia		
subjects affected / exposed	10 / 142 (7.04%)	3 / 141 (2.13%)
occurrences (all)	11	4
Hypomagnesaemia		
subjects affected / exposed	10 / 142 (7.04%)	11 / 141 (7.80%)
occurrences (all)	13	14

More information

Substantial protocol amendments (globally)

Were there any global substantial amendments to the protocol? Yes

Date	Amendment
30 January 2015	The amendment introduced the following changes: 1. The sample size reduction and power recalculation 2. A standardized definition for the assessment of NODM 3. The central reading of the tumor biopsy from the explanted liver of patients with HCC

Notes:

Interruptions (globally)

Were there any global interruptions to the trial? No

Limitations and caveats

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

Data recorded here are the CRAD001H2307 core study results. Updates with follow-up data for extension trial will be recorded in April 2019 (i.e., within 1 year of extension study completion).

Notes: