

**Clinical trial results:****A PHASE 2/3, RANDOMIZED, DOUBLE-MASKED, SHAM-CONTROLLED TRIAL OF QPI-1007 DELIVERED BY SINGLE OR MULTI-DOSE INTRAVITREAL INJECTION(S) TO SUBJECTS WITH ACUTE NONARTERITIC ANTERIOR ISCHEMIC OPTIC NEUROPATHY (NAION)****Summary**

EudraCT number	2015-003079-31
Trial protocol	DE IT
Global end of trial date	14 June 2019

Results information

Result version number	v1 (current)
This version publication date	22 July 2021
First version publication date	22 July 2021

Trial information**Trial identification**

Sponsor protocol code	QRK207
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Additional study identifiers

ISRCTN number	-
ClinicalTrials.gov id (NCT number)	NCT02341560
WHO universal trial number (UTN)	-

Notes:

Sponsors

Sponsor organisation name	Quark Pharmaceuticals, Inc.
Sponsor organisation address	7999 Gateway Boulevard, Suite 310, Newark, California , United States, 94560
Public contact	Clinical Operations, Quark Pharmaceuticals Inc., +1 510424020, dcafar@quarkpharma.com
Scientific contact	Clinical Operations, Quark Pharmaceuticals Inc., +1 510424020, dcafar@quarkpharma.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	14 June 2019
Is this the analysis of the primary completion data?	Yes
Primary completion date	14 June 2019
Global end of trial reached?	Yes
Global end of trial date	14 June 2019
Was the trial ended prematurely?	Yes

Notes:

General information about the trial

Main objective of the trial:

1. To assess the safety and tolerability of QPI-1007 IVT injections in subjects with recent-onset NAION.
2. To determine the effect on visual function of QPI-1007 IVT injections in subjects with recent-onset NAION.

Protection of trial subjects:

The trial was completed according to the International Conference on Harmonization (ICH) Guidelines for Good Clinical Practice. Compliance with this standard provides public assurance that the rights, safety, and well-being of trial subjects are protected, consistent with the principles that have their origin in the Declaration of Helsinki.

Background therapy: -

Evidence for comparator: -

Actual start date of recruitment	24 February 2016
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	Germany: 23
Country: Number of subjects enrolled	Italy: 23
Country: Number of subjects enrolled	Australia: 27
Country: Number of subjects enrolled	China: 34
Country: Number of subjects enrolled	India: 111
Country: Number of subjects enrolled	Israel: 47
Country: Number of subjects enrolled	Singapore: 4
Country: Number of subjects enrolled	United States: 463
Worldwide total number of subjects	732
EEA total number of subjects	46

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)	480
From 65 to 84 years	252
85 years and over	0

Subject disposition

Recruitment

Recruitment details:

The study was conducted at a total of 80 sites, with sites in Australia, China, Germany, India, Israel, Italy, Singapore and the United States

Pre-assignment

Screening details:

A total of 1,092 subjects with symptoms indicative of NAION were screened, of which 732 were randomized and 725 received test treatment within 16 days of onset of symptoms.

Period 1

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

Blinding implementation details:

Site study staff conducting and evaluating all ophthalmic evaluations will be masked to the treatment assignment, except those staff who perform the 30-minute post-dose assessments. Only masked staff should perform assessment of AEs for causality and relatedness to study treatment. Masked staff may not perform any Pharmacy tasks that compromise or could potentially compromise the study masking. The pharmacist, Investigator administering study treatment, unmasked coordinator will be unmasked.

Arms

Are arms mutually exclusive?	Yes
Arm title	Cohort 1 - ITT

Arm description:

Single dose regime, 1.5mg dose of QPI-1007 on day 1. Sham dose administered at months 2 and 4.

Arm type	Experimental
Investigational medicinal product name	QPI-1007
Investigational medicinal product code	
Other name	
Pharmaceutical forms	Solution for injection
Routes of administration	Intravitreal use

Dosage and administration details:

1.5 mg QPI-1007 dose. Preparation of QPI-1007 for administration should be done using aseptic techniques. The IVT injection procedure should be carried out under controlled aseptic conditions, which include the use of sterile gloves, a sterile drape, and a sterile eyelid speculum (or equivalent). Adequate topical anesthesia and a broad-spectrum topical microbicide should be given prior to the injection.

Arm title	Cohort 2 - ITT
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Arm description:

Single dose regime. 3mg QPI-1007 administered on day 1. Sham administration on months 2 and 4.

Arm type	Experimental
Investigational medicinal product name	QPI-1007
Investigational medicinal product code	
Other name	
Pharmaceutical forms	Solution for injection
Routes of administration	Intravitreal use

Dosage and administration details:

3mg QPI-1007 dose. Preparation of QPI-1007 for administration should be done using aseptic techniques. The IVT injection procedure should be carried out under controlled aseptic conditions, which include the use of sterile gloves, a sterile drape, and a sterile eyelid speculum (or equivalent). Adequate topical anesthesia and a broad-spectrum topical microbicide should be given prior to the injection.

Arm title	Cohort 3 - ITT
Arm description: Multiple dose regimen. 1.5 mg QPI-1007 dose on day 1, and months 2 and 4.	
Arm type	Experimental
Investigational medicinal product name	QPI-1007
Investigational medicinal product code	
Other name	
Pharmaceutical forms	Solution for injection
Routes of administration	Intravitreal use
Dosage and administration details: 1.5mg QPI-1007 dose. Preparation of QPI-1007 for administration should be done using aseptic techniques. The IVT injection procedure should be carried out under controlled aseptic conditions, which include the use of sterile gloves, a sterile drape, and a sterile eyelid speculum (or equivalent). Adequate topical anesthesia and a broad-spectrum topical microbicide should be given prior to the injection.	
Arm title	Cohort 4 - ITT
Arm description: Multiple dose regimen. 3mg QPI-1007 administered on day 1, and months 2 and 4.	
Arm type	Experimental
Investigational medicinal product name	QPI-1007
Investigational medicinal product code	
Other name	
Pharmaceutical forms	Solution for injection
Routes of administration	Intravitreal use
Dosage and administration details: 3mg QPI-1007 dose. Preparation of QPI-1007 for administration should be done using aseptic techniques. The IVT injection procedure should be carried out under controlled aseptic conditions, which include the use of sterile gloves, a sterile drape, and a sterile eyelid speculum (or equivalent). Adequate topical anesthesia and a broad-spectrum topical microbicide should be given prior to the injection.	
Arm title	Cohort 5 - ITT
Arm description: Sham control. For subjects receiving the sham-procedure no study drug was prepared but an empty syringe was used. The IVT injection was simulated by touching the eye with the blunt end of a syringe; the globe is not penetrated	
Arm type	No intervention
No investigational medicinal product assigned in this arm	

Number of subjects in period 1	Cohort 1 - ITT	Cohort 2 - ITT	Cohort 3 - ITT
Started	81	81	191
Completed	75	74	102
Not completed	6	7	89
Consent withdrawn by subject	1	4	6
Physician decision	-	-	1
Adverse event, non-fatal	1	1	2
Other	-	-	-
Study Terminated by Sponsor	-	-	75
Lost to follow-up	3	2	3

Lack of efficacy	1	-	-
Protocol deviation	-	-	2

Number of subjects in period 1	Cohort 4 - ITT	Cohort 5 - ITT
Started	188	191
Completed	102	103
Not completed	86	88
Consent withdrawn by subject	4	3
Physician decision	-	-
Adverse event, non-fatal	-	2
Other	2	-
Study Terminated by Sponsor	76	77
Lost to follow-up	1	2
Lack of efficacy	-	3
Protocol deviation	3	1

Baseline characteristics

Reporting groups

Reporting group title	Cohort 1 - ITT
Reporting group description: Single dose regime, 1.5mg dose of QPI-1007 on day 1. Sham dose administered at months 2 and 4.	
Reporting group title	Cohort 2 - ITT
Reporting group description: Single dose regime. 3mg QPI-1007 administered on day 1. Sham administration on months 2 and 4.	
Reporting group title	Cohort 3 - ITT
Reporting group description: Multiple dose regimen. 1.5 mg QPI-1007 dose on day 1, and months 2 and 4.	
Reporting group title	Cohort 4 - ITT
Reporting group description: Multiple dose regimen. 3mg QPI-1007 administered on day 1, and months 2 and 4.	
Reporting group title	Cohort 5 - ITT
Reporting group description: Sham control. For subjects receiving the sham-procedure no study drug was prepared but an empty syringe was used. The IVT injection was simulated by touching the eye with the blunt end of a syringe; the globe is not penetrated	

Reporting group values	Cohort 1 - ITT	Cohort 2 - ITT	Cohort 3 - ITT
Number of subjects	81	81	191
Age categorical Units: Subjects			
Adults (18-64 years)	53	60	124
From 65-84 years	28	21	67
Age continuous Units: years			
arithmetic mean	61.2	60.3	61.7
full range (min-max)	50 to 80	50 to 77	50 to 80
Gender categorical Units: Subjects			
Female	25	33	58
Male	56	48	133

Reporting group values	Cohort 4 - ITT	Cohort 5 - ITT	Total
Number of subjects	188	191	732
Age categorical Units: Subjects			
Adults (18-64 years)	122	121	480
From 65-84 years	66	70	252
Age continuous Units: years			
arithmetic mean	61.3	61.7	-
full range (min-max)	50 to 80	50 to 79	-

Gender categorical			
Units: Subjects			
Female	56	58	230
Male	132	133	502

Subject analysis sets

Subject analysis set title	Cohort 1 - mITT \leq 12days
Subject analysis set type	Modified intention-to-treat
Subject analysis set description: Single dose regime, 1.5mg dose of QPI-1007 on day 1. Sham dose administered at months 2 and 4. mITT \leq 12days analysis set include subjects who were randomized within 12 days of the onset of symptoms.	
Subject analysis set title	Cohort 2 - mITT \leq 12days
Subject analysis set type	Modified intention-to-treat
Subject analysis set description: Single dose regime. 3mg QPI-1007 administered on day 1. Sham administration on months 2 and 4. mITT \leq 12days analysis set include subjects who were randomized within 12 days of the onset of symptoms.	
Subject analysis set title	Cohort 3 - mITT \leq 12days
Subject analysis set type	Modified intention-to-treat
Subject analysis set description: Multiple dose regimen. 1.5 mg QPI-1007 dose on day 1, and months 2 and 4. mITT \leq 12days analysis set include subjects who were randomized within 12 days of the onset of symptoms.	
Subject analysis set title	Cohort 4 - mITT \leq 12days
Subject analysis set type	Modified intention-to-treat
Subject analysis set description: Multiple dose regimen. 3mg QPI-1007 administered on day 1, and months 2 and 4. mITT \leq 12days analysis set include subjects who were randomized within 12 days of the onset of symptoms.	
Subject analysis set title	Cohort 5 - mITT \leq 12days
Subject analysis set type	Modified intention-to-treat
Subject analysis set description: Sham control. mITT \leq 12days analysis set include subjects who were randomized within 12 days of the onset of symptoms.	
Subject analysis set title	Cohort 1 - As treated
Subject analysis set type	Per protocol
Subject analysis set description: Single dose regime, 1.5mg dose of QPI-1007 on day 1. Sham dose administered at months 2 and 4. As treated analysis set includes all subjects who received the correct test treatment.	
Subject analysis set title	Cohort 2 - As treated
Subject analysis set type	Per protocol
Subject analysis set description: Single dose regime. 3mg QPI-1007 administered on day 1. Sham administration on months 2 and 4. As treated analysis set includes all subjects who received the correct test treatment.	
Subject analysis set title	Cohort 3 - As treated
Subject analysis set type	Per protocol
Subject analysis set description: Multiple dose regimen. 1.5 mg QPI-1007 dose on day 1, and months 2 and 4. As treated analysis set includes all subjects who received the correct test treatment.	
Subject analysis set title	Cohort 4 - As treated
Subject analysis set type	Per protocol
Subject analysis set description: Multiple dose regimen. 3mg QPI-1007 administered on day 1, and months 2 and 4. As treated analysis set includes all subjects who received the correct test treatment.	
Subject analysis set title	Cohort 5 - As treated

Subject analysis set type	Per protocol
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Subject analysis set description:

Sham control. As treated analysis set includes all subjects who received the correct test treatment.

Reporting group values	Cohort 1 - mITT ≤ 12days	Cohort 2 - mITT ≤ 12days	Cohort 3 - mITT ≤ 12days
Number of subjects	101	100	168
Age categorical Units: Subjects			
Adults (18-64 years)	67	70	110
From 65-84 years	34	30	58
Age continuous Units: years arithmetic mean full range (min-max)			
Gender categorical Units: Subjects			
Female	29	38	54
Male	72	62	114

Reporting group values	Cohort 4 - mITT ≤ 12days	Cohort 5 - mITT ≤ 12days	Cohort 1 - As treated
Number of subjects	166	190	57
Age categorical Units: Subjects			
Adults (18-64 years)	111	120	
From 65-84 years	55	70	
Age continuous Units: years arithmetic mean full range (min-max)			61.3 50 to 80
Gender categorical Units: Subjects			
Female	49	58	16
Male	117	132	41

Reporting group values	Cohort 2 - As treated	Cohort 3 - As treated	Cohort 4 - As treated
Number of subjects	55	146	142
Age categorical Units: Subjects			
Adults (18-64 years)			
From 65-84 years			
Age continuous Units: years arithmetic mean full range (min-max)	61.0 50 to 80	61.5 50 to 80	60.9 50 to 80
Gender categorical Units: Subjects			
Female	18	46	41
Male	37	100	101

Reporting group values	Cohort 5 - As treated		

Number of subjects	144		
Age categorical			
Units: Subjects			
Adults (18-64 years)			
From 65-84 years			
Age continuous			
Units: years			
arithmetic mean	61.8		
full range (min-max)	50 to 79		
Gender categorical			
Units: Subjects			
Female	46		
Male	98		

End points

End points reporting groups

Reporting group title	Cohort 1 - ITT
Reporting group description: Single dose regime, 1.5mg dose of QPI-1007 on day 1. Sham dose administered at months 2 and 4.	
Reporting group title	Cohort 2 - ITT
Reporting group description: Single dose regime. 3mg QPI-1007 administered on day 1. Sham administration on months 2 and 4.	
Reporting group title	Cohort 3 - ITT
Reporting group description: Multiple dose regimen. 1.5 mg QPI-1007 dose on day 1, and months 2 and 4.	
Reporting group title	Cohort 4 - ITT
Reporting group description: Multiple dose regimen. 3mg QPI-1007 administered on day 1, and months 2 and 4.	
Reporting group title	Cohort 5 - ITT
Reporting group description: Sham control. For subjects receiving the sham-procedure no study drug was prepared but an empty syringe was used. The IVT injection was simulated by touching the eye with the blunt end of a syringe; the globe is not penetrated	
Subject analysis set title	Cohort 1 - mITT \leq 12days
Subject analysis set type	Modified intention-to-treat
Subject analysis set description: Single dose regime, 1.5mg dose of QPI-1007 on day 1. Sham dose administered at months 2 and 4. mITT \leq 12days analysis set include subjects who were randomized within 12 days of the onset of symptoms.	
Subject analysis set title	Cohort 2 - mITT \leq 12days
Subject analysis set type	Modified intention-to-treat
Subject analysis set description: Single dose regime. 3mg QPI-1007 administered on day 1. Sham administration on months 2 and 4. mITT \leq 12days analysis set include subjects who were randomized within 12 days of the onset of symptoms.	
Subject analysis set title	Cohort 3 - mITT \leq 12days
Subject analysis set type	Modified intention-to-treat
Subject analysis set description: Multiple dose regimen. 1.5 mg QPI-1007 dose on day 1, and months 2 and 4. mITT \leq 12days analysis set include subjects who were randomized within 12 days of the onset of symptoms.	
Subject analysis set title	Cohort 4 - mITT \leq 12days
Subject analysis set type	Modified intention-to-treat
Subject analysis set description: Multiple dose regimen. 3mg QPI-1007 administered on day 1, and months 2 and 4. mITT \leq 12days analysis set include subjects who were randomized within 12 days of the onset of symptoms.	
Subject analysis set title	Cohort 5 - mITT \leq 12days
Subject analysis set type	Modified intention-to-treat
Subject analysis set description: Sham control. mITT \leq 12days analysis set include subjects who were randomized within 12 days of the onset of symptoms.	
Subject analysis set title	Cohort 1 - As treated
Subject analysis set type	Per protocol
Subject analysis set description: Single dose regime, 1.5mg dose of QPI-1007 on day 1. Sham dose administered at months 2 and 4. As treated analysis set includes all subjects who received the correct test treatment.	
Subject analysis set title	Cohort 2 - As treated

Subject analysis set type	Per protocol
Subject analysis set description: Single dose regime. 3mg QPI-1007 administered on day 1. Sham administration on months 2 and 4. As treated analysis set includes all subjects who received the correct test treatment.	
Subject analysis set title	Cohort 3 - As treated
Subject analysis set type	Per protocol
Subject analysis set description: Multiple dose regimen. 1.5 mg QPI-1007 dose on day 1, and months 2 and 4. As treated analysis set includes all subjects who received the correct test treatment.	
Subject analysis set title	Cohort 4 - As treated
Subject analysis set type	Per protocol
Subject analysis set description: Multiple dose regimen. 3mg QPI-I007 administered on day 1, and months 2 and 4. As treated analysis set includes all subjects who received the correct test treatment.	
Subject analysis set title	Cohort 5 - As treated
Subject analysis set type	Per protocol
Subject analysis set description: Sham control. As treated analysis set includes all subjects who received the correct test treatment.	

Primary: Number of subjects who lose 15 letters or more in Best Corrected Visual Acuity (BCVA) score as measured by ETDRS visual acuity protocol in the study eye from Baseline to Month 6.

End point title	Number of subjects who lose 15 letters or more in Best Corrected Visual Acuity (BCVA) score as measured by ETDRS visual acuity protocol in the study eye from Baseline to Month 6.
End point description: Number of subjects who lose 15 letters or more in BCVA at Month 6 from Baseline. Results given as number of patients who have altered BCVA.	
End point type	Primary
End point timeframe: 6 months post treatment	

End point values	Cohort 1 - mITT ≤ 12days	Cohort 2 - mITT ≤ 12days	Cohort 3 - mITT ≤ 12days	Cohort 4 - mITT ≤ 12days
Subject group type	Subject analysis set	Subject analysis set	Subject analysis set	Subject analysis set
Number of subjects analysed	57	55	146	142
Units: Subjects	17	17	23	27

End point values	Cohort 5 - mITT ≤ 12days			
Subject group type	Subject analysis set			
Number of subjects analysed	144			
Units: Subjects	21			

Statistical analyses

Statistical analysis title	Percentage difference from control
Comparison groups	Cohort 1 - mITT \leq 12days v Cohort 5 - mITT \leq 12days
Number of subjects included in analysis	201
Analysis specification	Pre-specified
Analysis type	superiority
P-value	= 0.0217
Method	Regression, Logistic
Parameter estimate	Percentage difference
Confidence interval	
level	95 %
sides	2-sided

Statistical analysis title	Percentage difference from control
Comparison groups	Cohort 2 - mITT \leq 12days v Cohort 5 - mITT \leq 12days
Number of subjects included in analysis	199
Analysis specification	Pre-specified
Analysis type	superiority
P-value	= 0.0146
Method	Regression, Logistic
Parameter estimate	Percentage difference
Confidence interval	
level	95 %
sides	2-sided

Statistical analysis title	Percentage difference from control
Comparison groups	Cohort 3 - mITT \leq 12days v Cohort 5 - mITT \leq 12days
Number of subjects included in analysis	290
Analysis specification	Pre-specified
Analysis type	superiority
P-value	= 0.7808
Method	Regression, Logistic
Parameter estimate	Percentage difference
Confidence interval	
level	95 %
sides	2-sided

Statistical analysis title	Percentage difference from control
Comparison groups	Cohort 4 - mITT \leq 12days v Cohort 5 - mITT \leq 12days

Number of subjects included in analysis	286
Analysis specification	Pre-specified
Analysis type	superiority
P-value	= 0.293
Method	Regression, Logistic
Parameter estimate	Percentage difference
Confidence interval	
level	95 %
sides	2-sided

Adverse events

Adverse events information

Timeframe for reporting adverse events:

All AEs and SAEs will be recorded from the time of first study drug administration/sham procedure through 12 months post treatment or early termination

Assessment type	Systematic
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Dictionary used

Dictionary name	MedDRA
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Dictionary version	24.1
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Reporting groups

Reporting group title	Cohort 1 - As treated
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Reporting group description:

Single dose regime, 1.5mg dose of QPI-1007 on day 1. Sham dose administered at months 2 and 4. As treated analysis set includes all subjects who received the correct test treatment.

Reporting group title	Cohort 2 - As treated
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Reporting group description:

Single dose regime. 3mg QPI-1007 administered on day 1. Sham administration on months 2 and 4. As treated analysis set includes all subjects who received the correct test treatment.

Reporting group title	Cohort 3 - As treated
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Reporting group description:

Multiple dose regimen. 1.5 mg QPI-1007 dose on day 1, and months 2 and 4. As treated analysis set includes all subjects who received the correct test treatment.

Reporting group title	Cohort 4 - As treated
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Reporting group description:

Multiple dose regimen. 3mg QPI-1007 administered on day 1, and months 2 and 4. As treated analysis set includes all subjects who received the correct test treatment.

Reporting group title	Cohort 5 - As treated
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Reporting group description:

Sham control. As treated analysis set includes all subjects who received the correct test treatment.

Serious adverse events	Cohort 1 - As treated	Cohort 2 - As treated	Cohort 3 - As treated
Total subjects affected by serious adverse events			
subjects affected / exposed	11 / 101 (10.89%)	17 / 100 (17.00%)	17 / 168 (10.12%)
number of deaths (all causes)	0	0	1
number of deaths resulting from adverse events	0	0	1
Neoplasms benign, malignant and unspecified (incl cysts and polyps)			
Colon cancer			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Tumour haemorrhage			

subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Vascular disorders			
Additional description: Abdominal Aortic Aneurysm			
Aortic aneurysm			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Hypotension			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Hypertensive emergency			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
General disorders and administration site conditions			
Chest pain			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Multiple organ dysfunction syndrome			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Non-cardiac chest pain			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Respiratory, thoracic and mediastinal disorders			
Pulmonary embolism			

subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Psychiatric disorders			
Mental status changes	Additional description: Reaction to medication given for fall. reaction was altered mental state		
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Investigations			
C-Reactive Protein Increase			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Injury, poisoning and procedural complications			
Femoral neck fracture			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Postoperative ileus	Additional description: post operative		
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Cataract traumatic	Additional description: retraumatic cataract		
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Foreign body in eye	Additional description: foreign body in cornea		
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Contusion	Additional description: Hemorrhagic contusion		

subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Hip fracture	Additional description: fracture of left hip		
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Upper limb fracture			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Femur fracture			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Cardiac disorders			
Hypertensive crisis			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Atrial fibrillation			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Myocardial infarction	Additional description: non-ST segment elevation myocardial infarction		
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 2
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Ischaemic cardiomyopathy			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Ventricular fibrillation			

subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Coronary artery occlusion	Additional description: Chronic total occlusion of the coronary artery		
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Coronary artery disease			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Acute myocardial infarction			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Arteriosclerosis coronary artery			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Nervous system disorders			
Syncope			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Cerebral infarction			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	2 / 168 (1.19%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 2
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Cerebrovascular insufficiency			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Transient ischaemic attack			

subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Cerebrovascular accident			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Ischaemic stroke			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Subarachnoid haemorrhage			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Visual field defect			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Blood and lymphatic system disorders			
Anaemia	Additional description: worsening of anaemia		
subjects affected / exposed	1 / 101 (0.99%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Ear and labyrinth disorders			
Vertigo positional			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Vertigo	Additional description: Vertigo of unknown etiology		
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Eye disorders			

Vitreous floaters			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Macular hole	Additional description: Stage 4 macular hole		
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Age-related macular degeneration	Additional description: Wet age-related macular degeneration		
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Corneal perforation	Additional description: re-traumatic corneal perforation		
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Optic ischaemic neuropathy	Additional description: Progressive Anterior Optic Neuropathy Bilateral Sequential		
subjects affected / exposed	2 / 101 (1.98%)	2 / 100 (2.00%)	1 / 168 (0.60%)
occurrences causally related to treatment / all	0 / 2	0 / 2	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Blindness unilateral	Additional description: severe visual loss		
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Papilloedema	Additional description: Marked increase in Optic Nerve Edema OS due to NAION		
subjects affected / exposed	1 / 101 (0.99%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Photophobia	Additional description: Loss of vision to light perception OS		
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Visual impairment			

subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	1 / 168 (0.60%)
occurrences causally related to treatment / all	0 / 0	0 / 1	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Visual acuity reduced			
subjects affected / exposed	2 / 101 (1.98%)	8 / 100 (8.00%)	6 / 168 (3.57%)
occurrences causally related to treatment / all	0 / 2	0 / 8	0 / 6
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Optic atrophy			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Eye haemorrhage			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Gastrointestinal disorders			
Peptic ulcer			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Haemorrhoids			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Erosive oesophagitis			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Renal and urinary disorders			
Acute kidney injury			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	1 / 168 (0.60%)
occurrences causally related to treatment / all	0 / 0	0 / 1	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Musculoskeletal and connective tissue			

disorders	Additional description: Prolapsed Inter vertebral disc		
Intervertebral disc protrusion			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Infections and infestations			
Bronchitis viral			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 1	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Gastroenteritis viral			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Septic shock			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 1
Urinary tract infection			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Leptospirosis			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences causally related to treatment / all	0 / 0	0 / 0	0 / 1
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0
Metabolism and nutrition disorders			
Hyperkalaemia			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 1	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0	0 / 0

Serious adverse events	Cohort 4 - As treated	Cohort 5 - As treated	
Total subjects affected by serious adverse events			
subjects affected / exposed	22 / 166 (13.25%)	19 / 190 (10.00%)	

number of deaths (all causes)	0	0	
number of deaths resulting from adverse events	0	0	
Neoplasms benign, malignant and unspecified (incl cysts and polyps)			
Colon cancer			
subjects affected / exposed	1 / 166 (0.60%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Tumour haemorrhage			
subjects affected / exposed	0 / 166 (0.00%)	1 / 190 (0.53%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Vascular disorders			
Aortic aneurysm			
Additional description: Abdominal Aortic Aneurysm			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hypotension			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hypertensive emergency			
subjects affected / exposed	0 / 166 (0.00%)	1 / 190 (0.53%)	
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 pain			
subjects affected / exposed	0 / 166 (0.00%)	1 / 190 (0.53%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Multiple organ dysfunction syndrome			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Non-cardiac chest pain			

subjects affected / exposed	0 / 166 (0.00%)	1 / 190 (0.53%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Respiratory, thoracic and mediastinal disorders			
Pulmonary embolism			
subjects affected / exposed	0 / 166 (0.00%)	1 / 190 (0.53%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Psychiatric disorders			
Mental status changes			
Additional description: Reaction to medication given for fall. reaction was altered mental state			
subjects affected / exposed	0 / 166 (0.00%)	1 / 190 (0.53%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Investigations			
C-Reactive Protein Increase			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Injury, poisoning and procedural complications			
Femoral neck fracture			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Postoperative ileus			
Additional description: post operative			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Cataract traumatic			
Additional description: retraumatic cataract			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Foreign body in eye			
Additional description: foreign body in cornea			

subjects affected / exposed	1 / 166 (0.60%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Contusion	Additional description: Hemorrhagic contusion		
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hip fracture	Additional description: fracture of left hip		
subjects affected / exposed	0 / 166 (0.00%)	1 / 190 (0.53%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Upper limb fracture			
subjects affected / exposed	0 / 166 (0.00%)	1 / 190 (0.53%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Femur fracture			
subjects affected / exposed	1 / 166 (0.60%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Cardiac disorders			
Hypertensive crisis			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Atrial fibrillation			
subjects affected / exposed	1 / 166 (0.60%)	1 / 190 (0.53%)	
occurrences causally related to treatment / all	0 / 1	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Myocardial infarction	Additional description: non-ST segment elevation myocardial infarction		
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Ischaemic cardiomyopathy			

subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Ventricular fibrillation			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Coronary artery occlusion	Additional description: Chronic total occlusion of the coronary artery		
subjects affected / exposed	0 / 166 (0.00%)	1 / 190 (0.53%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Coronary artery disease			
subjects affected / exposed	1 / 166 (0.60%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Acute myocardial infarction			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Arteriosclerosis coronary artery			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Nervous system disorders			
Syncope			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Cerebral infarction			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Cerebrovascular insufficiency			

subjects affected / exposed	1 / 166 (0.60%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Transient ischaemic attack			
subjects affected / exposed	0 / 166 (0.00%)	1 / 190 (0.53%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Cerebrovascular accident			
subjects affected / exposed	1 / 166 (0.60%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Ischaemic stroke			
subjects affected / exposed	0 / 166 (0.00%)	1 / 190 (0.53%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Subarachnoid haemorrhage			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Visual field defect			
subjects affected / exposed	1 / 166 (0.60%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Blood and lymphatic system disorders			

Anaemia	Additional description: worsening of anaemia		
subjects affected / exposed	1 / 166 (0.60%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Ear and labyrinth disorders			
Vertigo positional			
subjects affected / exposed	1 / 166 (0.60%)	1 / 190 (0.53%)	
occurrences causally related to treatment / all	0 / 1	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	

Vertigo	Additional description: Vertigo of unknown etiology		

subjects affected / exposed	1 / 166 (0.60%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Eye disorders			
Vitreous floaters			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Macular hole			
Additional description: Stage 4 macular hole			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Age-related macular degeneration			
Additional description: Wet age-related macular degeneration			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Corneal perforation			
Additional description: re-traumatic corneal perforation			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Optic ischaemic neuropathy			
Additional description: Progressive Anterior Optic Neuropathy Bilateral Sequential			
subjects affected / exposed	4 / 166 (2.41%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 4	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Blindness unilateral			
Additional description: severe visual loss			
subjects affected / exposed	1 / 166 (0.60%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Papilloedema			
Additional description: Marked increase in Optic Nerve Edema OS due to NAION			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Photophobia			
Additional description: Loss of vision to light perception OS			

subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Visual impairment			
subjects affected / exposed	0 / 166 (0.00%)	2 / 190 (1.05%)	
occurrences causally related to treatment / all	0 / 0	0 / 2	
deaths causally related to treatment / all	0 / 0	0 / 0	
Visual acuity reduced			
subjects affected / exposed	7 / 166 (4.22%)	9 / 190 (4.74%)	
occurrences causally related to treatment / all	0 / 7	0 / 9	
deaths causally related to treatment / all	0 / 0	0 / 0	
Optic atrophy			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Eye haemorrhage			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Gastrointestinal disorders			
Peptic ulcer			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Haemorrhoids			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Erosive oesophagitis			
subjects affected / exposed	0 / 166 (0.00%)	1 / 190 (0.53%)	
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	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Musculoskeletal and connective tissue disorders			
Intervertebral disc protrusion	Additional description: Prolapsed Inter vertebral disc		
subjects affected / exposed	1 / 166 (0.60%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Infections and infestations			
Bronchitis viral			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Gastroenteritis viral			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Septic shock			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Urinary tract infection			
subjects affected / exposed	1 / 166 (0.60%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Leptospirosis			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences causally related to treatment / all	0 / 0	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Metabolism and nutrition disorders			
Hyperkalaemia			

subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)
occurrences causally related to treatment / all	0 / 0	0 / 0
deaths causally related to treatment / all	0 / 0	0 / 0

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

Non-serious adverse events	Cohort 1 - As treated	Cohort 2 - As treated	Cohort 3 - As treated
Total subjects affected by non-serious adverse events			
subjects affected / exposed	78 / 101 (77.23%)	75 / 100 (75.00%)	133 / 168 (79.17%)
Vascular disorders			
Arterial occlusive disease			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Hypertension			
subjects affected / exposed	3 / 101 (2.97%)	2 / 100 (2.00%)	7 / 168 (4.17%)
occurrences (all)	3	2	7
Hypotension			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Hyperaemia			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	0	0	0
Surgical and medical procedures			
Oral surgery			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences (all)	0	1	0
General disorders and administration site conditions			
Chest pain			
subjects affected / exposed	0 / 101 (0.00%)	2 / 100 (2.00%)	1 / 168 (0.60%)
occurrences (all)	0	2	1
Facial pain			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences (all)	0	1	0
Fatigue			

subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	1 / 100 (1.00%) 1	1 / 168 (0.60%) 1
Influenza like illness subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	1 / 168 (0.60%) 1
Glassy eyes subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	0 / 100 (0.00%) 0	0 / 168 (0.00%) 0
Injection site pain subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	1 / 168 (0.60%) 1
Injection site discomfort subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	0 / 100 (0.00%) 0	0 / 168 (0.00%) 0
Injection site erythema subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 2	0 / 100 (0.00%) 0	0 / 168 (0.00%) 0
Pain subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	0 / 100 (0.00%) 0	2 / 168 (1.19%) 2
Peripheral swelling subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	0 / 100 (0.00%) 0	2 / 168 (1.19%) 2
Pyrexia subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	4 / 100 (4.00%) 4	3 / 168 (1.79%) 3
Immune system disorders			
Drug hypersensitivity subjects affected / exposed occurrences (all)	2 / 101 (1.98%) 2	0 / 100 (0.00%) 0	0 / 168 (0.00%) 0
Hypersensitivity subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	0 / 168 (0.00%) 0
Seasonal allergy subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	1 / 168 (0.60%) 1

Bronchitis subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	1 / 100 (1.00%) 1	6 / 168 (3.57%) 6
Rhinitis subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	0 / 100 (0.00%) 0	0 / 168 (0.00%) 0
Social circumstances Menopause subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	0 / 100 (0.00%) 0	0 / 168 (0.00%) 0
Reproductive system and breast disorders Breast cyst subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	0 / 168 (0.00%) 0
Benign prostatic hyperplasia subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	0 / 100 (0.00%) 0	0 / 168 (0.00%) 0
Prostatitis subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	0 / 168 (0.00%) 0
Vaginal haemorrhage subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	0 / 168 (0.00%) 0
Respiratory, thoracic and mediastinal disorders Chronic obstructive pulmonary disease subjects affected / exposed occurrences (all)	2 / 101 (1.98%) 2	1 / 100 (1.00%) 1	0 / 168 (0.00%) 0
Cough subjects affected / exposed occurrences (all)	3 / 101 (2.97%) 3	2 / 100 (2.00%) 2	7 / 168 (4.17%) 7
Nasal congestion subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	0 / 100 (0.00%) 0	3 / 168 (1.79%) 3
Epistaxis			

subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences (all)	0	1	0
Hiccups			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences (all)	0	1	0
Nasal oedema			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Nasal polyps			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences (all)	1	0	1
Respiratory tract congestion			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Oropharyngeal pain			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	3 / 168 (1.79%)
occurrences (all)	0	0	3
Paranasal sinus discomfort			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences (all)	0	0	1
Rhinorrhoea			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	1 / 168 (0.60%)
occurrences (all)	0	1	1
Sleep apnoea syndrome			
subjects affected / exposed	6 / 101 (5.94%)	4 / 100 (4.00%)	3 / 168 (1.79%)
occurrences (all)	6	4	3
Asthma			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences (all)	0	1	0
Upper airway obstruction			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Upper respiratory tract congestion			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Psychiatric disorders			

Anxiety			
subjects affected / exposed	3 / 101 (2.97%)	2 / 100 (2.00%)	2 / 168 (1.19%)
occurrences (all)	3	2	2
Depression			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Insomnia			
subjects affected / exposed	1 / 101 (0.99%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences (all)	1	1	0
Stress			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Investigations			
Blood calcium increased			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences (all)	0	1	0
Blood cholesterol increased			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	2 / 168 (1.19%)
occurrences (all)	1	0	2
Blood homocysteine increased			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Blood pressure increased			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	1 / 168 (0.60%)
occurrences (all)	0	1	1
Blood urea increased			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	3 / 168 (1.79%)
occurrences (all)	0	0	3
Blood triglycerides increased			
subjects affected / exposed	4 / 101 (3.96%)	1 / 100 (1.00%)	1 / 168 (0.60%)
occurrences (all)	4	1	1
Injury, poisoning and procedural complications			
Toothache			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	4 / 168 (2.38%)
occurrences (all)	1	0	5
Tooth abscess			

subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Arthropod bite			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences (all)	0	1	0
Conjunctival abrasion			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Corneal abrasion			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences (all)	1	0	1
Contusion			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences (all)	0	1	0
Fall			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	1 / 168 (0.60%)
occurrences (all)	0	1	1
Foreign body in eye			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences (all)	0	0	1
Ligament sprain			
subjects affected / exposed	1 / 101 (0.99%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences (all)	1	1	0
Foot fracture			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Hand fracture			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences (all)	1	0	1
Injection related reaction			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Injury			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences (all)	0	1	0
Joint swelling			

subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	0	0	0
Joint injury			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	3 / 168 (1.79%)
occurrences (all)	0	0	3
Joint dislocation			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	0	0	0
Limb injury			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	1 / 168 (0.60%)
occurrences (all)	0	1	1
Muscle rupture			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	2 / 168 (1.19%)
occurrences (all)	0	0	2
Muscle strain			
subjects affected / exposed	2 / 101 (1.98%)	0 / 100 (0.00%)	2 / 168 (1.19%)
occurrences (all)	2	0	2
Lumbar vertebral fracture			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Post procedural haemorrhage			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	1 / 168 (0.60%)
occurrences (all)	0	1	1
Product administration error			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences (all)	1	0	1
Rib fracture			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences (all)	1	0	1
Stress fracture			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences (all)	0	1	0
Skin laceration			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Blood lactate dehydrogenase			

increased subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	0 / 100 (0.00%) 0	0 / 168 (0.00%) 0
Congenital, familial and genetic disorders Methylenetetrahydrofolate reductase gene mutation subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	0 / 168 (0.00%) 0
Cardiac disorders Atrial fibrillation subjects affected / exposed occurrences (all) Left ventricular hypertrophy subjects affected / exposed occurrences (all) Cardiomyopathy subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1 0 / 101 (0.00%) 0 1 / 101 (0.99%) 1	0 / 100 (0.00%) 0 1 / 100 (1.00%) 1 0 / 100 (0.00%) 0	0 / 168 (0.00%) 0 0 / 168 (0.00%) 0 0 / 168 (0.00%) 0
Nervous system disorders Visual field defect subjects affected / exposed occurrences (all) Nerve compression subjects affected / exposed occurrences (all)	3 / 101 (2.97%) 3 0 / 101 (0.00%) 0	3 / 100 (3.00%) 3 0 / 100 (0.00%) 0	4 / 168 (2.38%) 4 0 / 168 (0.00%) 0
Blood and lymphatic system disorders anaemia subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	2 / 100 (2.00%) 2	0 / 168 (0.00%) 0
Ear and labyrinth disorders Deafness subjects affected / exposed occurrences (all) Ear pain subjects affected / exposed occurrences (all) Tinnitus	1 / 101 (0.99%) 1 0 / 101 (0.00%) 0	0 / 100 (0.00%) 0 0 / 100 (0.00%) 0	0 / 168 (0.00%) 0 2 / 168 (1.19%) 2

subjects affected / exposed occurrences (all)	2 / 101 (1.98%) 2	0 / 100 (0.00%) 0	1 / 168 (0.60%) 1
Vertigo subjects affected / exposed occurrences (all)	2 / 101 (1.98%) 2	0 / 100 (0.00%) 0	0 / 168 (0.00%) 0
Eye disorders			
Abnormal sensation in eye subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	1 / 100 (1.00%) 1	0 / 168 (0.00%) 0
Anterior chamber cell subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	0 / 100 (0.00%) 0	0 / 168 (0.00%) 0
Blepharitis subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	2 / 100 (2.00%) 2	3 / 168 (1.79%) 3
Blepharospasm subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	0 / 100 (0.00%) 0	1 / 168 (0.60%) 1
Cataract subjects affected / exposed occurrences (all)	3 / 101 (2.97%) 3	2 / 100 (2.00%) 2	2 / 168 (1.19%) 2
Cataract nuclear subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	2 / 100 (2.00%) 2	1 / 168 (0.60%) 1
Cataract subcapsular subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	1 / 168 (0.60%) 1
Conjunctival haemorrhage subjects affected / exposed occurrences (all)	13 / 101 (12.87%) 13	11 / 100 (11.00%) 11	24 / 168 (14.29%) 24
Conjunctival hyperaemia subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	3 / 100 (3.00%) 3	3 / 168 (1.79%) 3
Corneal epithelial microcysts subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	0 / 168 (0.00%) 0

Cyanopsia			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences (all)	0	1	0
Dermatochalasis			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences (all)	0	1	0
Conjunctivitis allergic			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Cystoid macular oedema			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Diplopia			
subjects affected / exposed	1 / 101 (0.99%)	1 / 100 (1.00%)	2 / 168 (1.19%)
occurrences (all)	1	1	2
Dry eye			
subjects affected / exposed	3 / 101 (2.97%)	6 / 100 (6.00%)	7 / 168 (4.17%)
occurrences (all)	3	6	7
Eye disorder			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences (all)	0	1	0
Eye allergy			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Eye irritation			
subjects affected / exposed	2 / 101 (1.98%)	3 / 100 (3.00%)	4 / 168 (2.38%)
occurrences (all)	2	3	4
Eye pain			
subjects affected / exposed	5 / 101 (4.95%)	5 / 100 (5.00%)	19 / 168 (11.31%)
occurrences (all)	5	5	19
Eye swelling			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	2 / 168 (1.19%)
occurrences (all)	0	0	2
Eyelid oedema			
subjects affected / exposed	0 / 101 (0.00%)	2 / 100 (2.00%)	0 / 168 (0.00%)
occurrences (all)	0	2	0

Eyelid pain			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences (all)	0	1	0
Eye pruritus			
subjects affected / exposed	0 / 101 (0.00%)	2 / 100 (2.00%)	1 / 168 (0.60%)
occurrences (all)	0	2	1
swelling of the eyelid			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	0	0	0
Eyelid irritation			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Foreign body sensation in eyes			
subjects affected / exposed	2 / 101 (1.98%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	2	0	0
Glare			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	0	0	0
Macular fibrosis			
subjects affected / exposed	1 / 101 (0.99%)	2 / 100 (2.00%)	0 / 168 (0.00%)
occurrences (all)	1	2	0
Iritis			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences (all)	0	0	1
Lacrimation increased			
subjects affected / exposed	0 / 101 (0.00%)	2 / 100 (2.00%)	2 / 168 (1.19%)
occurrences (all)	0	2	2
Macular detachment			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences (all)	0	1	0
Macular oedema			
subjects affected / exposed	1 / 101 (0.99%)	3 / 100 (3.00%)	2 / 168 (1.19%)
occurrences (all)	1	3	2
Metamorphopsia			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	1 / 168 (0.60%)
occurrences (all)	0	1	1

Meibomian gland dysfunction subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	0 / 100 (0.00%) 0	1 / 168 (0.60%) 1
Ocular discomfort subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	1 / 100 (1.00%) 1	0 / 168 (0.00%) 0
Ocular hypertension subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	0 / 100 (0.00%) 0	1 / 168 (0.60%) 1
Ocular hyperaemia subjects affected / exposed occurrences (all)	2 / 101 (1.98%) 2	1 / 100 (1.00%) 1	3 / 168 (1.79%) 3
Optic atrophy subjects affected / exposed occurrences (all)	2 / 101 (1.98%) 2	2 / 100 (2.00%) 2	2 / 168 (1.19%) 2
Optic disc disorder subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	0 / 100 (0.00%) 0	1 / 168 (0.60%) 1
Optic disc haemorrhage subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	0 / 100 (0.00%) 0	5 / 168 (2.98%) 5
Optic ischaemic neuropathy subjects affected / exposed occurrences (all)	7 / 101 (6.93%) 7	4 / 100 (4.00%) 4	2 / 168 (1.19%) 2
Optic nerve disorder subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	1 / 168 (0.60%) 1
Optic nerve sheath haemorrhage subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	0 / 100 (0.00%) 0	2 / 168 (1.19%) 2
Papilloedema subjects affected / exposed occurrences (all)	6 / 101 (5.94%) 6	3 / 100 (3.00%) 3	3 / 168 (1.79%) 3
Photopsia subjects affected / exposed occurrences (all)	3 / 101 (2.97%) 3	2 / 100 (2.00%) 2	1 / 168 (0.60%) 1

Photophobia			
subjects affected / exposed	0 / 101 (0.00%)	3 / 100 (3.00%)	1 / 168 (0.60%)
occurrences (all)	0	3	1
Punctate keratitis			
subjects affected / exposed	1 / 101 (0.99%)	1 / 100 (1.00%)	6 / 168 (3.57%)
occurrences (all)	1	1	6
Pupillary reflex impaired			
subjects affected / exposed	2 / 101 (1.98%)	2 / 100 (2.00%)	1 / 168 (0.60%)
occurrences (all)	2	2	1
Retinal degeneration			
subjects affected / exposed	1 / 101 (0.99%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences (all)	1	1	0
Retinal disorder			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences (all)	0	0	1
Retinal detachment			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Retinal exudates			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	2 / 168 (1.19%)
occurrences (all)	1	0	2
Retinal haemorrhage			
subjects affected / exposed	1 / 101 (0.99%)	2 / 100 (2.00%)	2 / 168 (1.19%)
occurrences (all)	1	2	2
Retinal neovascularisation			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Retinoschisis			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences (all)	0	1	0
Vision blurred			
subjects affected / exposed	2 / 101 (1.98%)	1 / 100 (1.00%)	2 / 168 (1.19%)
occurrences (all)	2	1	2
Visual acuity reduced			
subjects affected / exposed	11 / 101 (10.89%)	18 / 100 (18.00%)	15 / 168 (8.93%)
occurrences (all)	11	18	15

Visual impairment subjects affected / exposed occurrences (all)	5 / 101 (4.95%) 5	6 / 100 (6.00%) 6	7 / 168 (4.17%) 7
Vitreous detachment subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	3 / 100 (3.00%) 3	2 / 168 (1.19%) 2
Vitreous floaters subjects affected / exposed occurrences (all)	5 / 101 (4.95%) 5	8 / 100 (8.00%) 8	5 / 168 (2.98%) 5
Vitritis subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	0 / 168 (0.00%) 0
Vitreous opacities subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	0 / 100 (0.00%) 0	1 / 168 (0.60%) 1
Asthenopia subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	0 / 100 (0.00%) 0	1 / 168 (0.60%) 1
Gastrointestinal disorders			
Diarrhoea subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	0 / 100 (0.00%) 0	1 / 168 (0.60%) 1
Abdominal pain subjects affected / exposed occurrences (all)	2 / 101 (1.98%) 2	3 / 100 (3.00%) 3	0 / 168 (0.00%) 0
Abdominal pain upper subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	0 / 100 (0.00%) 0	0 / 168 (0.00%) 0
Constipation subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	3 / 168 (1.79%) 3
Chronic gastritis subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	0 / 100 (0.00%) 0	0 / 168 (0.00%) 0
Dental caries			

subjects affected / exposed occurrences (all)	2 / 101 (1.98%) 2	0 / 100 (0.00%) 0	0 / 168 (0.00%) 0
Duodenal polyp subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	0 / 100 (0.00%) 0	0 / 168 (0.00%) 0
Inflammatory bowel disease subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	0 / 168 (0.00%) 0
Paraesthesia oral subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	0 / 168 (0.00%) 0
Rectal haemorrhage subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	0 / 168 (0.00%) 0
Nausea subjects affected / exposed occurrences (all)	2 / 101 (1.98%) 2	0 / 100 (0.00%) 0	1 / 168 (0.60%) 1
Vomiting subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	0 / 100 (0.00%) 0	0 / 168 (0.00%) 0
Hepatobiliary disorders Cholelithiasis subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	0 / 100 (0.00%) 0	0 / 168 (0.00%) 0
Skin and subcutaneous tissue disorders Alopecia subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	0 / 168 (0.00%) 0
Blister subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	1 / 168 (0.60%) 1
Dermatitis allergic subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	1 / 168 (0.60%) 1
Psoriasis			

subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences (all)	0	1	0
Eczema			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences (all)	0	1	0
Actinic keratosis			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Rash			
subjects affected / exposed	2 / 101 (1.98%)	1 / 100 (1.00%)	2 / 168 (1.19%)
occurrences (all)	2	1	2
Rash pruritic			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences (all)	0	1	0
Pruritus			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	0	0	0
Skin lesion			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences (all)	1	0	1
Impaired driving ability			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	1 / 168 (0.60%)
occurrences (all)	0	1	1
Splinter haemorrhages			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	0	0	0
Renal and urinary disorders			
Nephrolithiasis			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Endocrine disorders			
Hypothyroidism			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Diabetic retinopathy			

subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	1 / 100 (1.00%) 1	2 / 168 (1.19%) 2
Vitreous degeneration subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	0 / 168 (0.00%) 0
Infections and infestations			
Asymptomatic bacteriuria subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	0 / 100 (0.00%) 0	0 / 168 (0.00%) 0
Bronchiolitis subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	0 / 100 (0.00%) 0	0 / 168 (0.00%) 0
Cellulitis subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	1 / 100 (1.00%) 1	0 / 168 (0.00%) 0
Conjunctivitis subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	1 / 168 (0.60%) 1
Cystitis subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	0 / 100 (0.00%) 0	0 / 168 (0.00%) 0
Conjunctivitis viral subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	0 / 100 (0.00%) 0	1 / 168 (0.60%) 1
Eye infection subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	0 / 168 (0.00%) 0
Gastrointestinal infection subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	0 / 168 (0.00%) 0
Gastroenteritis viral subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	0 / 100 (0.00%) 0	1 / 168 (0.60%) 1
Helicobacter infection subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	0 / 100 (0.00%) 0	0 / 168 (0.00%) 0

Herpes zoster			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Hordeolum			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	1 / 168 (0.60%)
occurrences (all)	0	0	1
Influenza			
subjects affected / exposed	2 / 101 (1.98%)	2 / 100 (2.00%)	5 / 168 (2.98%)
occurrences (all)	2	2	5
Lower respiratory tract infection			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	1 / 168 (0.60%)
occurrences (all)	0	1	1
Labyrinthitis			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
lung infection			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Nail infection			
subjects affected / exposed	1 / 101 (0.99%)	0 / 100 (0.00%)	0 / 168 (0.00%)
occurrences (all)	1	0	0
Nasopharyngitis			
subjects affected / exposed	8 / 101 (7.92%)	11 / 100 (11.00%)	18 / 168 (10.71%)
occurrences (all)	8	11	18
Otitis externa			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	2 / 168 (1.19%)
occurrences (all)	0	0	2
Oral herpes			
subjects affected / exposed	0 / 101 (0.00%)	1 / 100 (1.00%)	0 / 168 (0.00%)
occurrences (all)	0	1	0
Pharyngitis			
subjects affected / exposed	0 / 101 (0.00%)	0 / 100 (0.00%)	2 / 168 (1.19%)
occurrences (all)	0	0	2
Pneumonia			
subjects affected / exposed	0 / 101 (0.00%)	2 / 100 (2.00%)	1 / 168 (0.60%)
occurrences (all)	0	2	1

Postoperative wound infection subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	0 / 168 (0.00%) 0
Root canal infection subjects affected / exposed occurrences (all)	1 / 101 (0.99%) 1	0 / 100 (0.00%) 0	0 / 168 (0.00%) 0
Sinusitis subjects affected / exposed occurrences (all)	6 / 101 (5.94%) 6	0 / 100 (0.00%) 0	9 / 168 (5.36%) 9
Tooth infection subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	1 / 168 (0.60%) 1
Upper respiratory tract infection subjects affected / exposed occurrences (all)	2 / 101 (1.98%) 2	0 / 100 (0.00%) 0	2 / 168 (1.19%) 2
Urinary tract infection subjects affected / exposed occurrences (all)	4 / 101 (3.96%) 4	1 / 100 (1.00%) 1	1 / 168 (0.60%) 1
Viral infection subjects affected / exposed occurrences (all)	2 / 101 (1.98%) 2	0 / 100 (0.00%) 0	0 / 168 (0.00%) 0
Vulval abscess subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	1 / 100 (1.00%) 1	0 / 168 (0.00%) 0
Chemical burns of eye subjects affected / exposed occurrences (all)	0 / 101 (0.00%) 0	0 / 100 (0.00%) 0	2 / 168 (1.19%) 2

Non-serious adverse events	Cohort 4 - As treated	Cohort 5 - As treated	
Total subjects affected by non-serious adverse events subjects affected / exposed	125 / 166 (75.30%)	134 / 190 (70.53%)	
Vascular disorders			
Arterial occlusive disease subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Hypertension			

subjects affected / exposed occurrences (all)	10 / 166 (6.02%) 10	10 / 190 (5.26%) 10	
Hypotension subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	0 / 190 (0.00%) 0	
Hyperaemia subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	0 / 190 (0.00%) 0	
Surgical and medical procedures Oral surgery subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
General disorders and administration site conditions Chest pain subjects affected / exposed occurrences (all)	2 / 166 (1.20%) 2	4 / 190 (2.11%) 4	
Facial pain subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Fatigue subjects affected / exposed occurrences (all)	2 / 166 (1.20%) 2	1 / 190 (0.53%) 1	
Influenza like illness subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	1 / 190 (0.53%) 1	
Glassy eyes subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Injection site pain subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	0 / 190 (0.00%) 0	
Injection site discomfort subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Injection site erythema			

subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	1 / 190 (0.53%) 1	
Pain subjects affected / exposed occurrences (all)	2 / 166 (1.20%) 2	1 / 190 (0.53%) 1	
Peripheral swelling subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	1 / 190 (0.53%) 1	
Pyrexia subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	2 / 190 (1.05%) 2	
Immune system disorders			
Drug hypersensitivity subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Hypersensitivity subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	1 / 190 (0.53%) 1	
Seasonal allergy subjects affected / exposed occurrences (all)	2 / 166 (1.20%) 2	5 / 190 (2.63%) 5	
Bronchitis subjects affected / exposed occurrences (all)	4 / 166 (2.41%) 4	2 / 190 (1.05%) 2	
Rhinitis subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Social circumstances			
Menopause subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Reproductive system and breast disorders			
Breast cyst subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Benign prostatic hyperplasia			

subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	1 / 190 (0.53%) 1	
Prostatitis subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Vaginal haemorrhage subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Respiratory, thoracic and mediastinal disorders			
Chronic obstructive pulmonary disease subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Cough subjects affected / exposed occurrences (all)	5 / 166 (3.01%) 5	4 / 190 (2.11%) 4	
Nasal congestion subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	2 / 190 (1.05%) 2	
Epistaxis subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	0 / 190 (0.00%) 0	
Hiccups subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Nasal oedema subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Nasal polyps subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Respiratory tract congestion subjects affected / exposed occurrences (all)	2 / 166 (1.20%) 1	0 / 190 (0.00%) 0	
Oropharyngeal pain			

subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	0 / 190 (0.00%) 0	
Paranasal sinus discomfort subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Rhinorrhoea subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	0 / 190 (0.00%) 0	
Sleep apnoea syndrome subjects affected / exposed occurrences (all)	5 / 166 (3.01%) 5	5 / 190 (2.63%) 5	
Asthma subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	1 / 190 (0.53%) 1	
Upper airway obstruction subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Upper respiratory tract congestion subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Psychiatric disorders			
Anxiety subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	2 / 190 (1.05%) 2	
Depression subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	3 / 190 (1.58%) 3	
Insomnia subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	0 / 190 (0.00%) 0	
Stress subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Investigations			
Blood calcium increased			

subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Blood cholesterol increased subjects affected / exposed occurrences (all)	2 / 166 (1.20%) 2	5 / 190 (2.63%) 5	
Blood homocysteine increased subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Blood pressure increased subjects affected / exposed occurrences (all)	2 / 166 (1.20%) 2	3 / 190 (1.58%) 3	
Blood urea increased subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	0 / 190 (0.00%) 0	
Blood triglycerides increased subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	3 / 190 (1.58%) 3	
Injury, poisoning and procedural complications			
Toothache subjects affected / exposed occurrences (all)	3 / 166 (1.81%) 3	1 / 190 (0.53%) 1	
Tooth abscess subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	1 / 190 (0.53%) 1	
Arthropod bite subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	1 / 190 (0.53%) 1	
Conjunctival abrasion subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Corneal abrasion subjects affected / exposed occurrences (all)	2 / 166 (1.20%) 2	1 / 190 (0.53%) 1	
Contusion			

subjects affected / exposed	2 / 166 (1.20%)	0 / 190 (0.00%)
occurrences (all)	2	0
Fall		
subjects affected / exposed	1 / 166 (0.60%)	3 / 190 (1.58%)
occurrences (all)	1	3
Foreign body in eye		
subjects affected / exposed	1 / 166 (0.60%)	2 / 190 (1.05%)
occurrences (all)	1	2
Ligament sprain		
subjects affected / exposed	0 / 166 (0.00%)	1 / 190 (0.53%)
occurrences (all)	0	1
Foot fracture		
subjects affected / exposed	1 / 166 (0.60%)	0 / 190 (0.00%)
occurrences (all)	1	0
Hand fracture		
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)
occurrences (all)	0	0
Injection related reaction		
subjects affected / exposed	1 / 166 (0.60%)	0 / 190 (0.00%)
occurrences (all)	1	0
Injury		
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)
occurrences (all)	0	0
Joint swelling		
subjects affected / exposed	2 / 166 (1.20%)	0 / 190 (0.00%)
occurrences (all)	2	0
Joint injury		
subjects affected / exposed	0 / 166 (0.00%)	1 / 190 (0.53%)
occurrences (all)	0	1
Joint dislocation		
subjects affected / exposed	2 / 166 (1.20%)	0 / 190 (0.00%)
occurrences (all)	2	0
Limb injury		
subjects affected / exposed	1 / 166 (0.60%)	0 / 190 (0.00%)
occurrences (all)	1	0
Muscle rupture		

subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Muscle strain subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	3 / 190 (1.58%) 3	
Lumbar vertebral fracture subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Post procedural haemorrhage subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Product administration error subjects affected / exposed occurrences (all)	2 / 166 (1.20%) 2	2 / 190 (1.05%) 2	
Rib fracture subjects affected / exposed occurrences (all)	2 / 166 (1.20%) 2	1 / 190 (0.53%) 1	
Stress fracture subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Skin laceration subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	3 / 190 (1.58%) 3	
Blood lactate dehydrogenase increased subjects affected / exposed occurrences (all)	2 / 166 (1.20%) 2	2 / 190 (1.05%) 2	
Congenital, familial and genetic disorders Methylenetetrahydrofolate reductase gene mutation subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	0 / 190 (0.00%) 0	
Cardiac disorders Atrial fibrillation subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	1 / 190 (0.53%) 1	
Left ventricular hypertrophy			

subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Cardiomyopathy subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Nervous system disorders Visual field defect subjects affected / exposed occurrences (all)	10 / 166 (6.02%) 10	4 / 190 (2.11%) 4	
Nerve compression subjects affected / exposed occurrences (all)	2 / 166 (1.20%) 2	0 / 190 (0.00%) 0	
Blood and lymphatic system disorders anaemia subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	1 / 190 (0.53%) 1	
Ear and labyrinth disorders Deafness subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Ear pain subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	0 / 190 (0.00%) 0	
Tinnitus subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	0 / 190 (0.00%) 0	
Vertigo subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Eye disorders Abnormal sensation in eye subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Anterior chamber cell subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Blepharitis			

subjects affected / exposed	2 / 166 (1.20%)	1 / 190 (0.53%)
occurrences (all)	2	1
Blepharospasm		
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)
occurrences (all)	0	0
Cataract		
subjects affected / exposed	3 / 166 (1.81%)	4 / 190 (2.11%)
occurrences (all)	3	4
Cataract nuclear		
subjects affected / exposed	1 / 166 (0.60%)	1 / 190 (0.53%)
occurrences (all)	1	1
Cataract subcapsular		
subjects affected / exposed	3 / 166 (1.81%)	0 / 190 (0.00%)
occurrences (all)	3	0
Conjunctival haemorrhage		
subjects affected / exposed	18 / 166 (10.84%)	4 / 190 (2.11%)
occurrences (all)	18	4
Conjunctival hyperaemia		
subjects affected / exposed	0 / 166 (0.00%)	4 / 190 (2.11%)
occurrences (all)	0	4
Corneal epithelial microcysts		
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)
occurrences (all)	0	0
Cyanopsia		
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)
occurrences (all)	0	0
Dermatochalasis		
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)
occurrences (all)	0	0
Conjunctivitis allergic		
subjects affected / exposed	0 / 166 (0.00%)	1 / 190 (0.53%)
occurrences (all)	0	1
Cystoid macular oedema		
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)
occurrences (all)	0	0
Diplopia		

subjects affected / exposed	1 / 166 (0.60%)	0 / 190 (0.00%)
occurrences (all)	1	0
Dry eye		
subjects affected / exposed	6 / 166 (3.61%)	6 / 190 (3.16%)
occurrences (all)	6	6
Eye disorder		
subjects affected / exposed	1 / 166 (0.60%)	0 / 190 (0.00%)
occurrences (all)	1	0
Eye allergy		
subjects affected / exposed	1 / 166 (0.60%)	2 / 190 (1.05%)
occurrences (all)	1	2
Eye irritation		
subjects affected / exposed	8 / 166 (4.82%)	1 / 190 (0.53%)
occurrences (all)	8	1
Eye pain		
subjects affected / exposed	8 / 166 (4.82%)	10 / 190 (5.26%)
occurrences (all)	8	10
Eye swelling		
subjects affected / exposed	1 / 166 (0.60%)	0 / 190 (0.00%)
occurrences (all)	1	0
Eyelid oedema		
subjects affected / exposed	0 / 166 (0.00%)	1 / 190 (0.53%)
occurrences (all)	0	1
Eyelid pain		
subjects affected / exposed	0 / 166 (0.00%)	10 / 190 (5.26%)
occurrences (all)	0	10
Eye pruritus		
subjects affected / exposed	2 / 166 (1.20%)	6 / 190 (3.16%)
occurrences (all)	2	6
swelling of the eyelid		
subjects affected / exposed	3 / 166 (1.81%)	0 / 190 (0.00%)
occurrences (all)	3	0
Eyelid irritation		
subjects affected / exposed	1 / 166 (0.60%)	0 / 190 (0.00%)
occurrences (all)	1	0
Foreign body sensation in eyes		

subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	2 / 190 (1.05%) 2
Glare subjects affected / exposed occurrences (all)	2 / 166 (1.20%) 2	0 / 190 (0.00%) 0
Macular fibrosis subjects affected / exposed occurrences (all)	3 / 166 (1.81%) 3	1 / 190 (0.53%) 1
Iritis subjects affected / exposed occurrences (all)	2 / 166 (1.20%) 2	1 / 190 (0.53%) 1
Lacrimation increased subjects affected / exposed occurrences (all)	4 / 166 (2.41%) 4	4 / 190 (2.11%) 4
Macular detachment subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0
Macular oedema subjects affected / exposed occurrences (all)	2 / 166 (1.20%) 2	1 / 190 (0.53%) 1
Metamorphopsia subjects affected / exposed occurrences (all)	2 / 166 (1.20%) 2	0 / 190 (0.00%) 0
Meibomian gland dysfunction subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	1 / 190 (0.53%) 1
Ocular discomfort subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	0 / 190 (0.00%) 0
Ocular hypertension subjects affected / exposed occurrences (all)	2 / 166 (1.20%) 2	0 / 190 (0.00%) 0
Ocular hyperaemia subjects affected / exposed occurrences (all)	3 / 166 (1.81%) 3	6 / 190 (3.16%) 6
Optic atrophy		

subjects affected / exposed	5 / 166 (3.01%)	1 / 190 (0.53%)
occurrences (all)	5	1
Optic disc disorder		
subjects affected / exposed	1 / 166 (0.60%)	2 / 190 (1.05%)
occurrences (all)	1	2
Optic disc haemorrhage		
subjects affected / exposed	3 / 166 (1.81%)	0 / 190 (0.00%)
occurrences (all)	3	0
Optic ischaemic neuropathy		
subjects affected / exposed	10 / 166 (6.02%)	8 / 190 (4.21%)
occurrences (all)	10	8
Optic nerve disorder		
subjects affected / exposed	1 / 166 (0.60%)	0 / 190 (0.00%)
occurrences (all)	1	0
Optic nerve sheath haemorrhage		
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)
occurrences (all)	0	0
Papilloedema		
subjects affected / exposed	9 / 166 (5.42%)	2 / 190 (1.05%)
occurrences (all)	9	2
Photopsia		
subjects affected / exposed	1 / 166 (0.60%)	1 / 190 (0.53%)
occurrences (all)	1	1
Photophobia		
subjects affected / exposed	4 / 166 (2.41%)	0 / 190 (0.00%)
occurrences (all)	4	0
Punctate keratitis		
subjects affected / exposed	4 / 166 (2.41%)	4 / 190 (2.11%)
occurrences (all)	4	4
Pupillary reflex impaired		
subjects affected / exposed	3 / 166 (1.81%)	1 / 190 (0.53%)
occurrences (all)	3	1
Retinal degeneration		
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)
occurrences (all)	0	0
Retinal disorder		

subjects affected / exposed	2 / 166 (1.20%)	2 / 190 (1.05%)
occurrences (all)	2	2
Retinal detachment		
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)
occurrences (all)	0	0
Retinal exudates		
subjects affected / exposed	2 / 166 (1.20%)	4 / 190 (2.11%)
occurrences (all)	2	4
Retinal haemorrhage		
subjects affected / exposed	1 / 166 (0.60%)	1 / 190 (0.53%)
occurrences (all)	1	1
Retinal neovascularisation		
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)
occurrences (all)	0	0
Retinoschisis		
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)
occurrences (all)	0	0
Vision blurred		
subjects affected / exposed	4 / 166 (2.41%)	5 / 190 (2.63%)
occurrences (all)	4	5
Visual acuity reduced		
subjects affected / exposed	21 / 166 (12.65%)	18 / 190 (9.47%)
occurrences (all)	21	18
Visual impairment		
subjects affected / exposed	3 / 166 (1.81%)	4 / 190 (2.11%)
occurrences (all)	3	4
Vitreous detachment		
subjects affected / exposed	8 / 166 (4.82%)	7 / 190 (3.68%)
occurrences (all)	8	7
Vitreous floaters		
subjects affected / exposed	5 / 166 (3.01%)	3 / 190 (1.58%)
occurrences (all)	5	3
Vitritis		
subjects affected / exposed	0 / 166 (0.00%)	1 / 190 (0.53%)
occurrences (all)	0	1
Vitreous opacities		

subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Asthenopia subjects affected / exposed occurrences (all)	2 / 166 (1.20%) 2	0 / 190 (0.00%) 0	
Gastrointestinal disorders			
Diarrhoea subjects affected / exposed occurrences (all)	3 / 166 (1.81%) 3	2 / 190 (1.05%) 2	
Abdominal pain subjects affected / exposed occurrences (all)	2 / 166 (1.20%) 2	1 / 190 (0.53%) 1	
Abdominal pain upper subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Constipation subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	1 / 190 (0.53%) 1	
Chronic gastritis subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Dental caries subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	0 / 190 (0.00%) 0	
Duodenal polyp subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Inflammatory bowel disease subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Paraesthesia oral subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Rectal haemorrhage subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	

Nausea			
subjects affected / exposed	1 / 166 (0.60%)	1 / 190 (0.53%)	
occurrences (all)	1	1	
Vomiting			
subjects affected / exposed	1 / 166 (0.60%)	2 / 190 (1.05%)	
occurrences (all)	1	2	
Hepatobiliary disorders			
Cholelithiasis			
subjects affected / exposed	1 / 166 (0.60%)	0 / 190 (0.00%)	
occurrences (all)	1	0	
Skin and subcutaneous tissue disorders			
Alopecia			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences (all)	0	0	
Blister			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences (all)	0	0	
Dermatitis allergic			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences (all)	0	0	
Psoriasis			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences (all)	0	0	
Eczema			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences (all)	0	0	
Actinic keratosis			
subjects affected / exposed	1 / 166 (0.60%)	0 / 190 (0.00%)	
occurrences (all)	1	0	
Rash			
subjects affected / exposed	2 / 166 (1.20%)	1 / 190 (0.53%)	
occurrences (all)	2	1	
Rash pruritic			
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)	
occurrences (all)	0	0	
Pruritus			

subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	0 / 190 (0.00%) 0	
Skin lesion subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Impaired driving ability subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Splinter haemorrhages subjects affected / exposed occurrences (all)	2 / 166 (1.20%) 2	1 / 190 (0.53%) 1	
Renal and urinary disorders Nephrolithiasis subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Endocrine disorders Hypothyroidism subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Diabetic retinopathy subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Vitreous degeneration subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	2 / 190 (1.05%) 2	
Infections and infestations Asymptomatic bacteriuria subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Bronchiolitis subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Cellulitis subjects affected / exposed occurrences (all)	1 / 166 (0.60%) 1	0 / 190 (0.00%) 0	
Conjunctivitis			

subjects affected / exposed	3 / 166 (1.81%)	3 / 190 (1.58%)
occurrences (all)	3	3
Cystitis		
subjects affected / exposed	2 / 166 (1.20%)	1 / 190 (0.53%)
occurrences (all)	2	1
Conjunctivitis viral		
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)
occurrences (all)	0	0
Eye infection		
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)
occurrences (all)	0	0
Gastrointestinal infection		
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)
occurrences (all)	0	0
Gastroenteritis viral		
subjects affected / exposed	1 / 166 (0.60%)	1 / 190 (0.53%)
occurrences (all)	1	1
Helicobacter infection		
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)
occurrences (all)	0	0
Herpes zoster		
subjects affected / exposed	1 / 166 (0.60%)	1 / 190 (0.53%)
occurrences (all)	1	1
Hordeolum		
subjects affected / exposed	2 / 166 (1.20%)	1 / 190 (0.53%)
occurrences (all)	2	1
Influenza		
subjects affected / exposed	2 / 166 (1.20%)	5 / 190 (2.63%)
occurrences (all)	2	5
Lower respiratory tract infection		
subjects affected / exposed	0 / 166 (0.00%)	1 / 190 (0.53%)
occurrences (all)	0	1
Labyrinthitis		
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)
occurrences (all)	0	0
lung infection		

subjects affected / exposed	1 / 166 (0.60%)	0 / 190 (0.00%)
occurrences (all)	1	0
Nail infection		
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)
occurrences (all)	0	0
Nasopharyngitis		
subjects affected / exposed	12 / 166 (7.23%)	17 / 190 (8.95%)
occurrences (all)	12	17
Otitis externa		
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)
occurrences (all)	0	0
Oral herpes		
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)
occurrences (all)	0	0
Pharyngitis		
subjects affected / exposed	1 / 166 (0.60%)	1 / 190 (0.53%)
occurrences (all)	1	1
Pneumonia		
subjects affected / exposed	0 / 166 (0.00%)	2 / 190 (1.05%)
occurrences (all)	0	2
Postoperative wound infection		
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)
occurrences (all)	0	0
Root canal infection		
subjects affected / exposed	0 / 166 (0.00%)	0 / 190 (0.00%)
occurrences (all)	0	0
Sinusitis		
subjects affected / exposed	5 / 166 (3.01%)	12 / 190 (6.32%)
occurrences (all)	5	12
Tooth infection		
subjects affected / exposed	0 / 166 (0.00%)	1 / 190 (0.53%)
occurrences (all)	0	1
Upper respiratory tract infection		
subjects affected / exposed	10 / 166 (6.02%)	5 / 190 (2.63%)
occurrences (all)	10	5
Urinary tract infection		

subjects affected / exposed occurrences (all)	4 / 166 (2.41%) 4	2 / 190 (1.05%) 2	
Viral infection subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	1 / 190 (0.53%) 1	
Vulval abscess subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	
Chemical burns of eye subjects affected / exposed occurrences (all)	0 / 166 (0.00%) 0	0 / 190 (0.00%) 0	

More information

Substantial protocol amendments (globally)

Were there any global substantial amendments to the protocol? Yes

Date	Amendment
13 January 2014	<p data-bbox="418 360 1126 389">Added language to note this is a Pivotal Phase II/III study</p> <p data-bbox="418 421 1390 535">Language added from provided reference regarding disease population. Revised confirmation of eligibility for subjects <50 and >80 years of age. Language modified to exclude subjects previously exposed to QPI-1007.</p> <p data-bbox="418 566 1050 595">Language added to note VF assessment at Month 2.</p> <p data-bbox="418 627 1353 680">Language added for AE and SAE reporting to cover local regulatory authority requirements for global protocol</p> <p data-bbox="418 712 1370 766">Language added for concomitant treatment collection to cover local regulatory authority requirements for global protocol.</p> <p data-bbox="418 797 1418 851">Added language to note that confirmation of enrollment eligibility to occur prior to Day 0.</p> <p data-bbox="418 882 858 911">Language added for SAE notification</p> <p data-bbox="418 943 1166 972">Optional interim analysis deleted due to revised study design.</p> <p data-bbox="418 1003 1401 1057">Language added to global protocol to capture video consent requirements where applicable, per local regulatory authorities.</p>

11 March 2015	<p>Clarified language to note dose options.</p> <p>Corrected language as not all subjects will receive three bi-monthly IVT injections.</p> <p>Interim analysis plan for the study was modified to include a futility analysis after 40 subjects in each treatment group complete Month 6 visit.</p> <p>Month 9 study visit added for safety follow-up.</p> <p>Language added to define afferent papillary defect inclusion criteria; Enrollment Adjudication Board removed and inclusion criteria confirmed that subjects must be 50 to 80 years of age.</p> <p>Added language to note Indian requirements, criterion for study eye, low dose aspirin therapy dose is per local standard. Added language to clarify exclusion criterion for diabetic retinopathy, concomitant disease and uveitis</p> <p>Language Added to allow basal cell carcinoma allowed only if adequately treated.</p> <p>Removal of FSH testing and ECG testing.</p> <p>Reference to diluted study drug removed as study drug will not be diluted.</p> <p>Correction of sample size determination and primary end point analysis, added multiple comparison procedure and futility analysis.</p> <p>Language added to note Key Secondary Endpoints and Secondary Efficacy Endpoints.</p> <p>Overall Summary of Related AEs table removed</p> <p>Contrast sensitivity added as an ophthalmologic assessment at Day 0 and Months 2, 4, 6, 9, and 12/ET.</p> <p>CRP and Westergren sedimentation rate added</p> <p>Entry criteria added for study eye, Instructions added for Day 0/Post-dose.</p> <p>Information added for adverse events in case of early treatment discontinuation</p> <p>Allocation of treatment groups defined for randomization</p> <p>Information on sensitivity and subgroup analysis was included.</p>
09 September 2015	<p>Changed all references for concomitant medications and treatments collection from 7 days to 14 days.</p> <p>Removed "differential" from WBC laboratory collection.</p> <p>Changed all references of Day "0" to Day "1" and Day "7" to Day "8."</p> <p>Added Europe to increase diversity of patient population.</p> <p>Adverse events, regardless of severity, must be collected for the entire study for all regions</p> <p>Added ganglion cell layer to Secondary objective.</p> <p>Study changed from 4-arm to 5-arm study. As such, study design was updated</p> <p>Reduced Screening window from 28 days to 14 days,</p> <p>Removed Contrast Sensitivity procedure/exam from protocol.</p> <p>Removed all mentions of 96 hours screening period and related references that stated Screening and Day 1 could occur on the same day.</p>

16 October 2015	<p>Updated: to reflect current understanding of disease process and number of subjects to accommodate recalculation of statistical hypothesis.</p> <p>Increased total number of study sites from 50 to 60.</p> <p>Removed aspirin requirement as background therapy.</p> <p>Added "OCT image and VF pattern compatible with the diagnosis of NAION, as determined by a Central Reading Center"</p> <p>image and VF pattern compatibility" to Screening visit task list.</p> <p>Changed inclusion criterion to allow Best-Corrected Visual Acuity score in the study eye to be better than or equal to 15 letter score without an upper limit</p> <p>Added exclusion criterion: "history of amiodarone use in the 12 months prior to the Day 1/Randomization visit." and "NAION secondary to acute blood loss' and "any intravitreal injection within 3 months prior to Day 1 in the study eye only:" and "diffuse pale swelling of the optic disc." and "bilateral (simultaneous) disc swelling</p> <p>Additional exclusion criteria added included: "Visual Field exclusions at Screening Visit (inconclusive visual fields should be sent to the Reading Center for adjudication)</p> <p>Updated study background to include new information and related references</p> <p>Added information regarding QRK208 study, 5th dosing cohort added in</p> <p>Changed "other efficacy endpoints."</p> <p>Allows for Screening and Day 1 to occur on same day.</p> <p>Change made to provide better understanding of per-visit procedures as not all procedures occur at all visits.</p>
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06 October 2016	<p>To explain purpose of and plan for newly-added sub-study.</p> <p>Clarification of global site distribution</p> <p>Clarification that there will be an earlier safety look, in addition to the already-listed interim analysis to be conducted after 40 subjects per group complete their Month 6 visit.</p> <p>It is not always the injecting physician who performs the 30-minute post-dose evaluations.</p> <p>Per German Regulatory Authority recommendation, removed barrier method as acceptable method of birth control</p> <p>Clarification of excluded ophthalmic surgeries.</p> <p>Clarification that subjects may be excluded by any visual field exclusions listed; subjects not required to meet ALL VF exclusions in order to be excluded. Clarified only study eye relevant to this criterion.</p> <p>Clarification that mild blepharitis, although considered an inflammatory condition, is NOT exclusionary.</p> <p>Correction to clarify potential subjects may be lactating and may not be actively nursing (lactating and nursing are not inclusive of one another).</p> <p>PDE5 inhibitors use is a risk for developing NAION and should not be used during the study.</p> <p>Updated to include new terms and correct existing list, to provide current information and status of QRK208 study.</p> <p>Spot size III using SITA standard 24-2 protocol is not done at Day 1, so this endpoint needs to be measured from Screening.</p> <p>Reduced requirement to avoid patient fatigue. Clarified spot size III requirement by name (previous version did not specify spot size).</p>
15 March 2018	<p>Sample size adjustment. The sample size has been recalculated to be consistent with the observed 20% overall rate of subjects losing 15-letters or more in visual acuity from 106 subjects to 213.</p> <p>Added language that another DMC will occur due to increased sample size</p> <p>Addition of ERG (for China sites only) to global protocol for Day 1 Pre-Dose, Month 2, and Month 6.</p> <p>Defined when 24-hour period starts (at first procedure after ICF, so any procedure after ICF).</p> <p>Added language found elsewhere in the protocol related to preferred time frame for study drug administration.</p> <p>Change type of stopper for study drug vial to "FuroTec-coated," instead of "Teflon-coated."</p> <p>Added statement that after the interim analysis, subjects already enrolled in the dropped regimen will continue until they have completed study participation</p> <p>Replaced duplicate IA information (already present in Sections 10.2 and 10.9) with definition of DMC purpose.</p>

Notes:

Interruptions (globally)

Were there any global interruptions to the trial? No

Limitations and caveats

None reported