

**Clinical trial results:**

A phase III long-term follow-up study to assess immunological memory induced following primary and booster vaccination with GSK Biologicals' 10-valent pneumococcal conjugate vaccine (10Pn-PD-DiT), through evaluation of early antibody responses after an additional dose of 10Pn-PD-DiT at 40-48 months of age, to evaluate in the fourth year of life, the immunogenicity and safety of a 2-dose catch-up immunization course with the 10Pn-PD-DiT vaccine and the impact of pneumococcal vaccination on nasopharyngeal carriage.

Summary

EudraCT number	2008-008104-41
Trial protocol	CZ
Global end of trial date	29 July 2011

Results information

Result version number	v1
This version publication date	20 June 2016
First version publication date	02 August 2015

Trial information**Trial identification**

Sponsor protocol code	112801
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Additional study identifiers

ISRCTN number	-
ClinicalTrials.gov id (NCT number)	NCT00950833
WHO universal trial number (UTN)	-

Notes:

Sponsors

Sponsor organisation name	GlaxoSmithKline Biologicals
Sponsor organisation address	Rue de l'Institut 89, Rixensart, Belgium, B-1330
Public contact	Clinical Trials Call Center, GlaxoSmithKline Biologicals, 044 2089-904466, GSKClinicalSupportHD@gsk.com
Scientific contact	Clinical Trials Call Center, GlaxoSmithKline Biologicals, 044 2089-904466, GSKClinicalSupportHD@gsk.com

Notes:

Paediatric regulatory details

Is trial part of an agreed paediatric investigation plan (PIP)	Yes
EMA paediatric investigation plan number(s)	EMA-000673-PIP01-09
Does article 45 of REGULATION (EC) No 1901/2006 apply to this trial?	No
Does article 46 of REGULATION (EC) No 1901/2006 apply to this trial?	Yes

Notes:

Results analysis stage

Analysis stage	Final
Date of interim/final analysis	19 August 2011
Is this the analysis of the primary completion data?	Yes
Primary completion date	16 September 2010
Global end of trial reached?	Yes
Global end of trial date	29 July 2011
Was the trial ended prematurely?	No

Notes:

General information about the trial

Main objective of the trial:

•To demonstrate the immunological memory induced following primary vaccination in study 10PN-PD-DIT-010 (107017) and booster vaccination in study 10PN-PD-DIT-014 (107137) with the 10Pn-PD-DiT vaccine, through evaluation of early antibody responses after an additional dose of 10Pn-PD-DiT at 40-48 months of age, compared to the unprimed group.

Criteria for immune memory:

The immune memory will be demonstrated if the lower limit of the 95% CI around the GMC ratios (pooled primed groups over unprimed group) is higher than 1 for all 10 vaccine serotypes.

Protection of trial subjects:

The vaccines were observed closely for at least 30 minutes following the administration of vaccine(s), with appropriate medical treatment readily available in case of a rare anaphylactic reaction.

Background therapy: -

Evidence for comparator: -

Actual start date of recruitment	10 August 2009
Long term follow-up planned	No
Independent data monitoring committee (IDMC) involvement?	No

Notes:

Population of trial subjects

Subjects enrolled per country

Country: Number of subjects enrolled	Czech Republic: 466
Worldwide total number of subjects	466
EEA total number of subjects	466

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)	466
Adolescents (12-17 years)	0
Adults (18-64 years)	0
From 65 to 84 years	0
85 years and over	0

Subject disposition

Recruitment

Recruitment details: -

Pre-assignment

Screening details:

During the screening the following steps occurred: check for inclusion/exclusion criteria, contraindications/precautions, medical history of the subjects and signing informed consent forms.

Pre-assignment period milestones

Number of subjects started	466
Number of subjects completed	443

Pre-assignment subject non-completion reasons

Reason: Number of subjects	Vaccine not administered: 5
Reason: Number of subjects	Consent withdrawn by subject: 18

Period 1

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

Arms

Are arms mutually exclusive?	Yes
Arm title	10Pn-PD-DiT/Paracetamol Group

Arm description:

Subjects were vaccinated with three primary vaccination doses with prophylactic administration of paracetamol in study 10PN-PD-DIT-010 (107017), and a booster dose of GSK Biologicals' Synflorix™ vaccine, co-administered with Infanrix™ hexa and with prophylactic antipyretic treatment (rectal paracetamol) in study 10PN-PD-DIT-014 BST-010 and who received in this study one dose of Synflorix™ vaccine at month 9 (40-48 months of age).

Arm type	Experimental
Investigational medicinal product name	10-valent pneumococcal conjugate vaccine
Investigational medicinal product code	
Other name	10Pn, 10Pn-PD-DiT, GlaxoSmithKline (GSK) Biologicals' 10-valent pneumococcal conjugate vaccine, Synflorix™, GlaxoSmithKline (GSK) Biologicals' 1024850A vaccine
Pharmaceutical forms	Suspension for injection
Routes of administration	Intramuscular use

Dosage and administration details:

One dose administered intramuscularly in the deltoid muscle at Month 9 (40-48 month of age).

Arm title	10Pn-PD-DiT Group
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Arm description:

Subjects were vaccinated with three primary vaccination doses with prophylactic administration of paracetamol in study 10PN-PD-DIT-010 (107017), at 12-15 months of age a booster dose of GSK Biologicals' Synflorix™ vaccine, co-administered with Infanrix™ hexa without prophylactic antipyretic treatment (rectal paracetamol) in study 10PN-PD-DIT-014 BST-010 (107137) and one dose of Synflorix™ vaccine in this study at Month 9 (month 40-48 of age).

Arm type	Experimental
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Investigational medicinal product name	10-valent Streptococcus pneumoniae conjugate vaccine
Investigational medicinal product code	
Other name	10Pn, 10Pn-PD-DiT, GlaxoSmithKline (GSK) Biologicals' 10-valent pneumococcal conjugate vaccine, Synflorix™, GlaxoSmithKline (GSK) Biologicals' 1024850A vaccine
Pharmaceutical forms	Suspension for injection
Routes of administration	Intramuscular use

Dosage and administration details:

One dose administered intramuscularly in the deltoid region at Month 9 (month 40-48 of age).

Arm title	Unprimed Group
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Arm description:

Subjects were vaccinated with three primary vaccination doses without prophylactic administration of paracetamol in study 10PN-PD-DIT-010 (107017), at 12-15 months of age a booster dose of GSK Biologicals' 10Pn-PD-DiT vaccine, co-administered with Infanrix™ hexa without prophylactic antipyretic treatment (rectal) in study 10PN-PD-DIT-014 BST-010 (107137) and two doses, one of the 10Pn-PD-DiT vaccine at Month 9 (40-48 months of age) and one at Month 11 (42-50 months of age).

Arm type	Active comparator
Investigational medicinal product name	10-valent Streptococcus pneumoniae conjugate vaccine
Investigational medicinal product code	
Other name	10Pn, 10Pn-PD-DiT, GlaxoSmithKline (GSK) Biologicals' 10-valent pneumococcal conjugate vaccine, Synflorix™, GlaxoSmithKline (GSK) Biologicals' 1024850A vaccine
Pharmaceutical forms	Suspension for injection
Routes of administration	Intramuscular use

Dosage and administration details:

Two doses administered intramuscularly in the deltoid muscle at Month 9 (40-48 months of age) and Month 11 (42-50 months of age).

Number of subjects in period 1^[1]	10Pn-PD-DiT/Paracetamol Group	10Pn-PD-DiT Group	Unprimed Group
Started	112	108	223
Completed	112	108	223

Notes:

[1] - The number of subjects reported to be in the baseline period are not the same as the worldwide number enrolled in the trial. It is expected that these numbers will be the same.

Justification: 5 of the enrolled subjects were allocated subject numbers but did not receive vaccination, hence they were excluded from the study. Also, 18 subjects dropped out of the study prior to start and were eliminated from the Total vaccinated cohort.

Baseline characteristics

Reporting groups

Reporting group title	10Pn-PD-DiT/Paracetamol Group
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Reporting group description:

Subjects were vaccinated with three primary vaccination doses with prophylactic administration of paracetamol in study 10PN-PD-DIT-010 (107017), and a booster dose of GSK Biologicals' Synflorix™ vaccine, co-administered with Infanrix™ hexa and with prophylactic antipyretic treatment (rectal paracetamol) in study 10PN-PD-DIT-014 BST-010 and who received in this study one dose of Synflorix™ vaccine at month 9 (40-48 months of age).

Reporting group title	10Pn-PD-DiT Group
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Reporting group description:

Subjects were vaccinated with three primary vaccination doses with prophylactic administration of paracetamol in study 10PN-PD-DIT-010 (107017), at 12-15 months of age a booster dose of GSK Biologicals' Synflorix™ vaccine, co-administered with Infanrix™ hexa without prophylactic antipyretic treatment (rectal paracetamol) in study 10PN-PD-DIT-014 BST-010 (107137) and one dose of Synflorix™ vaccine in this study at Month 9 (month 40-48 of age).

Reporting group title	Unprimed Group
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Reporting group description:

Subjects were vaccinated with three primary vaccination doses without prophylactic administration of paracetamol in study 10PN-PD-DIT-010 (107017), at 12-15 months of age a booster dose of GSK Biologicals' 10Pn-PD-DiT vaccine, co-administered with Infanrix™ hexa without prophylactic antipyretic treatment (rectal) in study 10PN-PD-DIT-014 BST-010 (107137) and two doses, one of the 10Pn-PD-DiT vaccine at Month 9 (40-48 months of age) and one at Month 11 (42-50 months of age).

Reporting group values	10Pn-PD-DiT/Paracetamol Group	10Pn-PD-DiT Group	Unprimed Group
Number of subjects	112	108	223
Age categorical Units: Subjects			
In utero Preterm newborn infants (gestational age < 37 wks) Newborns (0-27 days) Infants and toddlers (28 days-23 months) Children (2-11 years) Adolescents (12-17 years) Adults (18-64 years) From 65-84 years 85 years and over			
Age continuous Units: months			
arithmetic mean	39.2	39.1	37.7
standard deviation	± 1.53	± 1.54	± 3.36
Gender categorical Units: Subjects			
Female	60	48	99
Male	52	60	124

Reporting group values	Total		
Number of subjects	443		

Age categorical Units: Subjects			
In utero	0		
Preterm newborn infants (gestational age < 37 wks)	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)	0		
From 65-84 years	0		
85 years and over	0		
Age continuous Units: months			
arithmetic mean			
standard deviation	-		
Gender categorical Units: Subjects			
Female	207		
Male	236		

Subject analysis sets

Subject analysis set title	Poled primed Group
Subject analysis set type	Sub-group analysis

Subject analysis set description:

AP-AP and NAP-pre Groups were pooled into Primed Group.

Reporting group values	Poled primed Group		
Number of subjects	220		
Age categorical Units: Subjects			
In utero			
Preterm newborn infants (gestational age < 37 wks)			
Newborns (0-27 days)			
Infants and toddlers (28 days-23 months)			
Children (2-11 years)			
Adolescents (12-17 years)			
Adults (18-64 years)			
From 65-84 years			
85 years and over			
Age continuous Units: months			
arithmetic mean	39.15		
standard deviation	± 1.53		
Gender categorical Units: Subjects			
Female	108		
Male	112		

End points

End points reporting groups

Reporting group title	10Pn-PD-DiT/Paracetamol Group
Reporting group description: Subjects were vaccinated with three primary vaccination doses with prophylactic administration of paracetamol in study 10PN-PD-DIT-010 (107017), and a booster dose of GSK Biologicals' Synflorix™ vaccine, co-administered with Infanrix™ hexa and with prophylactic antipyretic treatment (rectal paracetamol) in study 10PN-PD-DIT-014 BST-010 and who received in this study one dose of Synflorix™ vaccine at month 9 (40-48 months of age).	
Reporting group title	10Pn-PD-DiT Group
Reporting group description: Subjects were vaccinated with three primary vaccination doses with prophylactic administration of paracetamol in study 10PN-PD-DIT-010 (107017), at 12-15 months of age a booster dose of GSK Biologicals' Synflorix™ vaccine, co-administered with Infanrix™ hexa without prophylactic antipyretic treatment (rectal paracetamol) in study 10PN-PD-DIT-014 BST-010 (107137) and one dose of Synflorix™ vaccine in this study at Month 9 (month 40-48 of age).	
Reporting group title	Unprimed Group
Reporting group description: Subjects were vaccinated with three primary vaccination doses without prophylactic administration of paracetamol in study 10PN-PD-DIT-010 (107017), at 12-15 months of age a booster dose of GSK Biologicals' 10Pn-PD-DiT vaccine, co-administered with Infanrix™ hexa without prophylactic antipyretic treatment (rectal) in study 10PN-PD-DIT-014 BST-010 (107137) and two doses, one of the 10Pn-PD-DiT vaccine at Month 9 (40-48 months of age) and one at Month 11 (42-50 months of age).	
Subject analysis set title	Poled primed Group
Subject analysis set type	Sub-group analysis
Subject analysis set description: AP-AP and NAP-pre Groups were pooled into Primed Group.	

Primary: Antibody concentrations against pneumococcal serotypes 1, 4, 5, 6B, 7F, 9V, 14, 18C, 19F and 23F (Anti-1, 4, 5, 6B, 7F, 9V, 14, 18C, 19F and 23F).

End point title	Antibody concentrations against pneumococcal serotypes 1, 4, 5, 6B, 7F, 9V, 14, 18C, 19F and 23F (Anti-1, 4, 5, 6B, 7F, 9V, 14, 18C, 19F and 23F). ^[1]
End point description:	
End point type	Primary
End point timeframe: At 7-10 days after the single/first dose of 10-valent pneumococcal conjugate vaccine.	

Notes:

[1] - The end point is not reporting statistics for all the arms in the baseline period. It is expected all the baseline period arms will be reported on when providing values for an end point on the baseline period.
Justification: This analysis was based on the pooled primed group that included the subjects in the remaining baseline groups.

End point values	Unprimed Group	Poled primed Group		
Subject group type	Reporting group	Subject analysis set		
Number of subjects analysed	204	215		
Units: µg/mL				
geometric mean (confidence interval 95%)				
Anti-1 (N=215; 204)	1.24 (1.07 to 1.42)	7.63 (6.57 to 8.86)		
Anti-4 (N=215; 204)	4.52 (3.96 to 5.17)	12.95 (11.37 to 14.75)		

Anti-5 (N=215; 202)	0.72 (0.63 to 0.84)	9.76 (8.44 to 11.29)		
Anti-6B (N=215; 203)	0.27 (0.22 to 0.33)	7.67 (6.7 to 8.78)		
Anti-7F (N=215; 204)	1.37 (1.19 to 1.58)	6.51 (5.69 to 7.46)		
Anti-9V (N=213; 202)	0.69 (0.58 to 0.83)	9.75 (8.46 to 11.24)		
Anti-14 (N=214; 204)	1.01 (0.8 to 1.27)	23.07 (20.03 to 26.57)		
Anti-18C (N=214; 204)	3.25 (2.71 to 3.9)	32.54 (28.15 to 37.61)		
Anti-19F (N=214;204)	4.31 (3.59 to 5.17)	39.84 (34.12 to 46.51)		
Anti-23F (N=215; 204)	0.25 (0.2 to 0.32)	9.24 (7.86 to 10.86)		

Statistical analyses

Statistical analysis title	Immune response non-inferiority - Anti-1
Statistical analysis description:	
To demonstrate the immunological memory induced following primary vaccination in study 10PN-PD-DIT-010 (107017) and booster vaccination in study 10PN-PD-DIT-014 (107137) with the 10Pn-PD-DiT vaccine, through evaluation of early antibody responses after an additional dose of 10Pn-PD-DiT vaccine at 40-48 months of age, compared to the Unprimed Group.	
Comparison groups	Unprimed Group v Poled primed Group
Number of subjects included in analysis	419
Analysis specification	Pre-specified
Analysis type	non-inferiority ^[2]
Parameter estimate	GMC ratio
Point estimate	6.17
Confidence interval	
level	95 %
sides	2-sided
lower limit	5.03
upper limit	7.58

Notes:

[2] - The immune memory would be demonstrated if the lower limit of the 95% confidence interval (CI) around the geometric mean concentration (GMC) ratios (Pooled Primed Group over Unprimed Group) was higher than 1 for all 10 vaccine serotypes.

Statistical analysis title	Immune response non-inferiority - Anti-4
Statistical analysis description:	
To demonstrate the immunological memory induced following primary vaccination in study 10PN-PD-DIT-010 (107017) and booster vaccination in study 10PN-PD-DIT-014 (107137) with the 10Pn-PD-DiT vaccine, through evaluation of early antibody responses after an additional dose of 10Pn-PD-DiT vaccine at 40-48 months of age, compared to the Unprimed Group.	
Comparison groups	Unprimed Group v Poled primed Group
Number of subjects included in analysis	419
Analysis specification	Pre-specified
Analysis type	non-inferiority ^[3]
Parameter estimate	GMC ratio
Point estimate	2.86

Confidence interval	
level	95 %
sides	2-sided
lower limit	2.38
upper limit	3.45

Notes:

[3] - The immune memory would be demonstrated if the lower limit of the 95% confidence interval (CI) around the geometric mean concentration (GMC) ratios (Pooled Primed Group over Unprimed Group) was higher than 1 for all 10 vaccine serotypes.

Statistical analysis title	Immune response non-inferiority - Anti-5
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Statistical analysis description:

To demonstrate the immunological memory induced following primary vaccination in study 10PN-PD-DIT-010 (107017) and booster vaccination in study 10PN-PD-DIT-014 (107137) with the 10Pn-PD-DiT vaccine, through evaluation of early antibody responses after an additional dose of 10Pn-PD-DiT vaccine at 40-48 months of age, compared to the Unprimed Group.

Comparison groups	Unprimed Group v Poled primed Group
Number of subjects included in analysis	419
Analysis specification	Pre-specified
Analysis type	non-inferiority ^[4]
Parameter estimate	GMC ratio
Point estimate	13.47
Confidence interval	
level	95 %
sides	2-sided
lower limit	10.96
upper limit	16.55

Notes:

[4] - The immune memory would be demonstrated if the lower limit of the 95% confidence interval (CI) around the geometric mean concentration (GMC) ratios (Pooled Primed Group over Unprimed Group) was higher than 1 for all 10 vaccine serotypes.

Statistical analysis title	Immune response non-inferiority - Anti-6B
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Statistical analysis description:

To demonstrate the immunological memory induced following primary vaccination in study 10PN-PD-DIT-010 (107017) and booster vaccination in study 10PN-PD-DIT-014 (107137) with the 10Pn-PD-DiT vaccine, through evaluation of early antibody responses after an additional dose of 10Pn-PD-DiT vaccine at 40-48 months of age, compared to the Unprimed Group.

Comparison groups	Unprimed Group v Poled primed Group
Number of subjects included in analysis	419
Analysis specification	Pre-specified
Analysis type	non-inferiority ^[5]
Parameter estimate	GMC ratio
Point estimate	28.81
Confidence interval	
level	95 %
sides	2-sided
lower limit	22.54
upper limit	36.81

Notes:

[5] - The immune memory would be demonstrated if the lower limit of the 95% confidence interval (CI) around the geometric mean concentration (GMC) ratios (Pooled Primed Group over Unprimed Group) was higher than 1 for all 10 vaccine serotypes.

Statistical analysis title	Immune response non-inferiority - Anti-7F
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Statistical analysis description:

To demonstrate the immunological memory induced following primary vaccination in study 10PN-PD-DIT-010 (107017) and booster vaccination in study 10PN-PD-DIT-014 (107137) with the 10Pn-PD-DiT vaccine, through evaluation of early antibody responses after an additional dose of 10Pn-PD-DiT vaccine at 40-48 months of age, compared to the Unprimed Group.

Comparison groups	Unprimed Group v Poled primed Group
Number of subjects included in analysis	419
Analysis specification	Pre-specified
Analysis type	non-inferiority ^[6]
Parameter estimate	GMC ratio
Point estimate	4.75
Confidence interval	
level	95 %
sides	2-sided
lower limit	3.9
upper limit	5.78

Notes:

[6] - The immune memory would be demonstrated if the lower limit of the 95% confidence interval (CI) around the geometric mean concentration (GMC) ratios (Pooled Primed Group over Unprimed Group) was higher than 1 for all 10 vaccine serotypes.

Statistical analysis title	Immune response non-inferiority - Anti-9V
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Statistical analysis description:

To demonstrate the immunological memory induced following primary vaccination in study 10PN-PD-DIT-010 (107017) and booster vaccination in study 10PN-PD-DIT-014 (107137) with the 10Pn-PD-DiT vaccine, through evaluation of early antibody responses after an additional dose of 10Pn-PD-DiT vaccine at 40-48 months of age, compared to the Unprimed Group.

Comparison groups	Unprimed Group v Poled primed Group
Number of subjects included in analysis	419
Analysis specification	Pre-specified
Analysis type	non-inferiority ^[7]
Parameter estimate	GMC ratio
Point estimate	14.1
Confidence interval	
level	95 %
sides	2-sided
lower limit	11.21
upper limit	17.75

Notes:

[7] - The immune memory would be demonstrated if the lower limit of the 95% confidence interval (CI) around the geometric mean concentration (GMC) ratios (Pooled Primed Group over Unprimed Group) was higher than 1 for all 10 vaccine serotypes.

Statistical analysis title	Immune response non-inferiority - Anti-14
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Statistical analysis description:

To demonstrate the immunological memory induced following primary vaccination in study 10PN-PD-DIT-010 (107017) and booster vaccination in study 10PN-PD-DIT-014 (107137) with the 10Pn-PD-DiT vaccine, through evaluation of early antibody responses after an additional dose of 10Pn-PD-DiT vaccine at 40-48 months of age, compared to the Unprimed Group.

Comparison groups	Unprimed Group v Poled primed Group
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Number of subjects included in analysis	419
Analysis specification	Pre-specified
Analysis type	non-inferiority ^[8]
Parameter estimate	GMC ratio
Point estimate	22.92
Confidence interval	
level	95 %
sides	2-sided
lower limit	17.51
upper limit	30

Notes:

[8] - The immune memory would be demonstrated if the lower limit of the 95% confidence interval (CI) around the geometric mean concentration (GMC) ratios (Pooled Primed Group over Unprimed Group) was higher than 1 for all 10 vaccine serotypes.

Statistical analysis title	Immune response non-inferiority - Anti-18C
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Statistical analysis description:

To demonstrate the immunological memory induced following primary vaccination in study 10PN-PD-DIT-010 (107017) and booster vaccination in study 10PN-PD-DIT-014 (107137) with the 10Pn-PD-DiT vaccine, through evaluation of early antibody responses after an additional dose of 10Pn-PD-DiT vaccine at 40-48 months of age, compared to the Unprimed Group.

Comparison groups	Unprimed Group v Poled primed Group
Number of subjects included in analysis	419
Analysis specification	Pre-specified
Analysis type	non-inferiority ^[9]
Parameter estimate	GMC ratio
Point estimate	10.01
Confidence interval	
level	95 %
sides	2-sided
lower limit	7.95
upper limit	12.61

Notes:

[9] - The immune memory would be demonstrated if the lower limit of the 95% confidence interval (CI) around the geometric mean concentration (GMC) ratios (Pooled Primed Group over Unprimed Group) was higher than 1 for all 10 vaccine serotypes.

Statistical analysis title	Immune response non-inferiority - Anti-19F
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Statistical analysis description:

To demonstrate the immunological memory induced following primary vaccination in study 10PN-PD-DIT-010 (107017) and booster vaccination in study 10PN-PD-DIT-014 (107137) with the 10Pn-PD-DiT vaccine, through evaluation of early antibody responses after an additional dose of 10Pn-PD-DiT vaccine at 40-48 months of age, compared to the Unprimed Group.

Comparison groups	Unprimed Group v Poled primed Group
Number of subjects included in analysis	419
Analysis specification	Pre-specified
Analysis type	non-inferiority ^[10]
Parameter estimate	GMC ratio
Point estimate	9.25
Confidence interval	
level	95 %
sides	2-sided
lower limit	7.29
upper limit	11.74

Notes:

[10] - The immune memory would be demonstrated if the lower limit of the 95% confidence interval (CI) around the geometric mean concentration (GMC) ratios (Pooled Primed Group over Unprimed Group) was higher than 1 for all 10 vaccine serotypes.

Statistical analysis title	Immune response non-inferiority - Anti-23F
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Statistical analysis description:

To demonstrate the immunological memory induced following primary vaccination in study 10PN-PD-DIT-010 (107017) and booster vaccination in study 10PN-PD-DIT-014 (107137) with the 10Pn-PD-DiT vaccine, through evaluation of early antibody responses after an additional dose of 10Pn-PD-DiT vaccine at 40-48 months of age, compared to the Unprimed Group.

Comparison groups	Unprimed Group v Poled primed Group
Number of subjects included in analysis	419
Analysis specification	Pre-specified
Analysis type	non-inferiority ^[11]
Parameter estimate	GMC ratio
Point estimate	36.52
Confidence interval	
level	95 %
sides	2-sided
lower limit	27.59
upper limit	48.34

Notes:

[11] - The immune memory would be demonstrated if the lower limit of the 95% confidence interval (CI) around the geometric mean concentration (GMC) ratios (Pooled Primed Group over Unprimed Group) was higher than 1 for all 10 vaccine serotypes.

Secondary: Anti-1, anti-4, anti-5, anti-6B, anti-7F, anti-9V, anti-14, anti-18C, anti 19F and anti23-F antibody concentrations.

End point title	Anti-1, anti-4, anti-5, anti-6B, anti-7F, anti-9V, anti-14, anti-18C, anti 19F and anti23-F antibody concentrations. ^[12]
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End point description:

End point type	Secondary
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End point timeframe:

Prior to the single/first dose of 10-valent pneumococcal conjugate vaccine.

Notes:

[12] - The end point is not reporting statistics for all the arms in the baseline period. It is expected all the baseline period arms will be reported on when providing values for an end point on the baseline period.

Justification: This analysis was based on the pooled primed group that included the subjects in the remaining baseline groups.

End point values	Unprimed Group	Poled primed Group		
Subject group type	Reporting group	Subject analysis set		
Number of subjects analysed	209	215		
Units: µg/mL				
geometric mean (confidence interval 95%)				
Anti-1 (N=215; 208)	0.09 (0.08 to 0.11)	0.27 (0.23 to 0.33)		
Anti-4 (N=213; 206)	0.05 (0.04 to 0.06)	0.2 (0.17 to 0.23)		
Anti-5 (N=212; 208)	0.1 (0.09 to 0.11)	0.41 (0.36 to 0.47)		
Anti-6B (N=212; 202)	0.1 (0.08 to 0.12)	0.8 (0.61 to 1.05)		

Anti-7F (N=215; 209)	0.06 (0.05 to 0.07)	0.48 (0.41 to 0.56)		
Anti-9V (N=211; 203)	0.07 (0.06 to 0.09)	0.5 (0.41 to 0.61)		
Anti-14 (N=213; 203)	0.28 (0.22 to 0.36)	1.21 (0.97 to 1.5)		
Anti-18C (N=215; 205)	0.09 (0.07 to 0.11)	0.65 (0.54 to 0.79)		
Anti-19F (N=215; 208)	0.44 (0.32 to 0.59)	2.35 (1.75 to 3.15)		
Anti-23F (N=215; 207)	0.08 (0.07 to 0.1)	0.96 (0.72 to 1.29)		

Statistical analyses

No statistical analyses for this end point

Secondary: Opsonophagocytic activity (OPA) titres against pneumococcal serotypes 1, 4, 5, 6B, 7F, 9V, 14, 18C, 19F and 23F.

End point title	Opsonophagocytic activity (OPA) titres against pneumococcal serotypes 1, 4, 5, 6B, 7F, 9V, 14, 18C, 19F and 23F. ^[13]
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End point description:

End point type	Secondary
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End point timeframe:

Prior to (PRE) and 7-10 days after (POST) the single/first dose of 10-valent pneumococcal conjugate vaccine.

Notes:

[13] - The end point is not reporting statistics for all the arms in the baseline period. It is expected all the baseline period arms will be reported on when providing values for an end point on the baseline period.

Justification: This analysis was based on the pooled primed group that included the subjects in the remaining baseline groups.

End point values	Unprimed Group	Poled primed Group		
Subject group type	Reporting group	Subject analysis set		
Number of subjects analysed	191	199		
Units: Titres				
geometric mean (confidence interval 95%)				
Opsono-1, PRE (N=199; 191)	5.4 (4.7 to 6.2)	10.2 (8 to 12.9)		
Opsono-1, POST (N=194; 182)	632 (524.4 to 761.7)	3106.5 (2670 to 3614.5)		
Opsono-4, PRE (N=189; 174)	9.1 (6.8 to 12.1)	18.1 (13.1 to 24.9)		
Opsono-4, POST (N=195; 187)	13109.9 (11080.6 to 15510.7)	27273.3 (22682.9 to 32792.6)		
Opsono-5, PRE (N=199; 185)	4 (4 to 4.1)	8.9 (7.6 to 10.5)		
Opsono-5, POST (N=194; 182)	145.8 (111.7 to 190.3)	1020 (874 to 1190.2)		
Opsono-6B, PRE (N=183; 173)	46.7 (29.8 to 73.3)	163.5 (108.5 to 246.4)		

Opsono-6B, POST (N=192; 184)	1472.2 (1053.2 to 2057.8)	5789.5 (4637.3 to 7227.9)		
Opsono-7F, PRE (N=187; 167)	973.4 (798.5 to 1186.6)	1112.3 (951.4 to 1300.4)		
Opsono-7F, POST (N=191; 186)	13647.4 (11801.9 to 15781.3)	19988.9 (16834.5 to 23734.2)		
Opsono-9V, PRE (N=171; 156)	268.2 (192.2 to 374.3)	481.8 (394.6 to 588.1)		
Opsono-9V, POST (N=194; 186)	14668.8 (12476.1 to 17247)	17952.5 (14699.2 to 21925.7)		
Opsono-14, PRE (N=184; 166)	145.3 (97.5 to 216.6)	267.6 (196.9 to 363.8)		
Opsono-14, POST (N=195; 187)	4454.3 (3921.1 to 5059.9)	16256.8 (13573 to 19471.4)		
Opsono-18C, PRE (N=186; 169)	5.7 (4.7 to 7)	14.2 (10.6 to 19.2)		
Opsono-18C, POST (N=188; 180)	9092.2 (7381.2 to 11199.9)	7413.8 (6072.3 to 9051.6)		
Opsono-19F, PRE (N=192; 186)	12.9 (9.6 to 17.2)	79.5 (55.4 to 113.9)		
Opsono-19F, POST (N=192; 190)	902.5 (659.2 to 1235.6)	6271.1 (4814.8 to 8168)		
Opsono-23F, PRE (N=189; 166)	220.6 (132 to 368.7)	462.7 (292.9 to 730.9)		
Opsono-23F, POST (N=197; 189)	5776.5 (4686.8 to 7119.5)	15613.5 (12386.9 to 19680.6)		

Statistical analyses

No statistical analyses for this end point

Secondary: Antibody concentrations against pneumococcal serotypes 6A and 19A (anti-6A and 19A).

End point title	Antibody concentrations against pneumococcal serotypes 6A and 19A (anti-6A and 19A). ^[14]
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End point description:

End point type	Secondary
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End point timeframe:

Prior to and 7-10 days after the single/first dose of 10-valent pneumococcal conjugate vaccine.

Notes:

[14] - The end point is not reporting statistics for all the arms in the baseline period. It is expected all the baseline period arms will be reported on when providing values for an end point on the baseline period.

Justification: This analysis was based on the pooled primed group that included the subjects in the remaining baseline groups.

End point values	Unprimed Group	Poled primed Group		
Subject group type	Reporting group	Subject analysis set		
Number of subjects analysed	207	215		
Units: µg/mL				
geometric mean (confidence interval 95%)				
Anti-6A, PRE (N=214; 207)	0.11 (0.09 to 0.13)	0.39 (0.3 to 0.5)		
Anti-6A, PI (N=215; 203)	0.2 (0.17 to 0.25)	2.4 (2.01 to 2.85)		
Anti-19A, PRE (N=215; 207)	0.22 (0.18 to 0.28)	0.52 (0.41 to 0.67)		
Anti-19A, PI (N=214; 203)	0.65 (0.52 to 0.82)	6.75 (5.41 to 8.41)		

Statistical analyses

No statistical analyses for this end point

Secondary: Opsonophagocytic activity (OPA) titres against pneumococcal cross-reactive serotypes 6A and 19A.

End point title	Opsonophagocytic activity (OPA) titres against pneumococcal cross-reactive serotypes 6A and 19A. ^[15]
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End point description:

End point type	Secondary
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End point timeframe:

Prior to and 7-10 days after the single/first dose of 10-valent pneumococcal conjugate vaccine.

Notes:

[15] - The end point is not reporting statistics for all the arms in the baseline period. It is expected all the baseline period arms will be reported on when providing values for an end point on the baseline period.

Justification: This analysis was based on the pooled primed group that included the subjects in the remaining baseline groups.

End point values	Unprimed Group	Poled primed Group		
Subject group type	Reporting group	Subject analysis set		
Number of subjects analysed	183	191		
Units: Titres				
geometric mean (confidence interval 95%)				
Opsono-6A, PRE (N=178; 167)	49 (32.4 to 74)	95.2 (65.3 to 138.7)		
Opsono-6A, PI (N=191; 180)	863.3 (619.3 to 1203.4)	2408.4 (1815.7 to 3194.5)		
Opsono-19A, PRE (N=190; 183)	880.5 (622.7 to 1245)	15.9 (11.7 to 21.6)		
Opsono-19A, PI (N=191; 175)	597.6 (467.2 to 764.4)	2104.9 (1560.9 to 2838.7)		

Statistical analyses

No statistical analyses for this end point

Secondary: Concentrations of antibodies against protein D (anti-PD).

End point title	Concentrations of antibodies against protein D (anti-PD). ^[16]
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End point description:

End point type	Secondary
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End point timeframe:

Prior to and 7-10 days after the single/first dose of 10-valent pneumococcal conjugate vaccine.

Notes:

[16] - The end point is not reporting statistics for all the arms in the baseline period. It is expected all the baseline period arms will be reported on when providing values for an end point on the baseline period.

Justification: This analysis was based on the pooled primed group that included the subjects in the remaining baseline groups.

End point values	Unprimed Group	Poled primed Group		
Subject group type	Reporting group	Subject analysis set		
Number of subjects analysed	204	215		
Units: EL.U/mL				
geometric mean (confidence interval 95%)				
Anti-PD, PRE (N=210; 198)	105.6 (93.8 to 119)	464.2 (401.6 to 536.5)		
Anti-PD, PI (N=215; 204)	374.3 (315.1 to 444.6)	2673.7 (2359.1 to 3030.2)		

Statistical analyses

No statistical analyses for this end point

Secondary: Memory B-cell detection for serotype specific polysaccharides (PS) (polysaccharides 1, 5, 6B, 18C, 19F, 23F and C).

End point title	Memory B-cell detection for serotype specific polysaccharides (PS) (polysaccharides 1, 5, 6B, 18C, 19F, 23F and C).
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End point description:

End point type	Secondary
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End point timeframe:

Prior to (PRE) and 7-10 days (POST) after the single/first dose of 10-valent pneumococcal conjugate vaccine.

End point values	10Pn-PD-DiT/Paracetamol Group	10Pn-PD-DiT Group	Unprimed Group	
Subject group type	Reporting group	Reporting group	Reporting group	
Number of subjects analysed	112	107	210	
Units: mIU/ml				
arithmetic mean (standard deviation)				
1 PS, PRE	288.6 (± 352.5)	269.5 (± 432.2)	254.8 (± 368.4)	81.5 (± 109.2)
5 PS, PRE	139.4 (± 214.7)	141.9 (± 232.4)		
6 PS, PRE	372 (± 545.5)	327.2 (± 554.4)	639.9 (± 846.5)	
18C PS, PRE	537.7 (± 641.8)	530.9 (± 693.8)	135.2 (± 200.6)	
19F PS, PRE	169.7 (± 415.3)	149.1 (± 194.4)	164.5 (± 343.4)	
23F PS, PRE	112.6 (± 176.1)	285 (± 484.3)	185.6 (± 270.1)	
C-PS, PRE	475.9 (± 682.3)	462.5 (± 704.5)	469.9 (± 639.2)	
1 PS, POST	1488.8 (± 1584.2)	1017.4 (± 1309.9)	755.9 (± 1017.4)	
5 PS, POST	233.5 (± 235.7)	406.6 (± 447.8)	152.6 (± 231)	
6 PS, POST	629.1 (± 773.2)	1265.8 (± 1738.6)	526 (± 648.6)	
18C PS, POST	3839 (± 5960.5)	8308.4 (± 7166.9)	2053.8 (± 2122.2)	
19F PS, POST	1056.7 (± 1334)	1092.4 (± 1724.3)	708.2 (± 1609.4)	
23F PS, POST	579.8 (± 632.5)	1123.7 (± 2095.4)	327.9 (± 470.5)	
C-PS, POST	679.2 (± 885.8)	915.5 (± 1191.3)	762.6 (± 1039.4)	

Statistical analyses

No statistical analyses for this end point

Secondary: Anti-1, anti-4, anti-5, anti-6B, anti-7F, anti-9V, anti-14, anti-18C, anti 19F and anti23-F antibody concentrations.

End point title	Anti-1, anti-4, anti-5, anti-6B, anti-7F, anti-9V, anti-14, anti-18C, anti 19F and anti23-F antibody concentrations. ^[17]
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End point description:

End point type	Secondary
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End point timeframe:

1 month after vaccination with the second dose of 10-valent pneumococcal conjugate vaccine (Month 12).

Notes:

[17] - The end point is not reporting statistics for all the arms in the baseline period. It is expected all the baseline period arms will be reported on when providing values for an end point on the baseline period.

Justification: This analysis was based on the unprimed group only.

End point values	Unprimed Group			
Subject group type	Reporting group			
Number of subjects analysed	209			
Units: µg/mL				
geometric mean (confidence interval 95%)				
Anti-1 (N=209)	2.33 (2.13 to 2.54)			
Anti-4 (N=209)	6.26 (5.73 to 6.84)			
Anti-5 (N=209)	2.69 (2.44 to 2.96)			
Anti-6B (N=209)	0.84 (0.73 to 0.97)			
Anti-7F (N=208)	3.63 (3.33 to 3.95)			
Anti-9V (N=209)	1.73 (1.56 to 1.93)			
Anti-14 (N=209)	5.21 (4.6 to 5.89)			
Anti-18C (N=209)	13.59 (11.91 to 15.51)			
Anti-19F (N=209)	11.83 (10.35 to 13.52)			
Anti-23F (N=209)	0.99 (0.86 to 1.15)			

Statistical analyses

No statistical analyses for this end point

Secondary: Opsono-1, opsono-4, opsono-5, opsono-6B, opsono-7F, opsono-9V, opsoo-14, opsono-18C, opsono-19F and opsono-23F titres.

End point title	Opsono-1, opsono-4, opsono-5, opsono-6B, opsono-7F, opsono-9V, opsoo-14, opsono-18C, opsono-19F and opsono-23F titres. ^[18]
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End point description:

End point type	Secondary
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End point timeframe:

1 month after vaccination with the second dose of 10-valent pneumococcal conjugate vaccine (Month 12).

Notes:

[18] - The end point is not reporting statistics for all the arms in the baseline period. It is expected all the baseline period arms will be reported on when providing values for an end point on the baseline period.

Justification: This analysis was based on the unprimed group only.

End point values	Unprimed Group			
Subject group type	Reporting group			
Number of subjects analysed	201			
Units: Tires				
geometric mean (confidence interval 95%)				
Opsono-1 (N=201)	125.6 (105 to 150.1)			
Opsono-4 (N=200)	2451.2 (2203.6 to 2726.8)			
Opsono-5 (N=198)	57.2 (47.3 to 69.1)			
Opsono-6B (N=196)	1345 (1066.8 to 1695.7)			
Opsono-7F (N=198)	6527.2 (5836.5 to 7299.6)			
Opsono-9V (N=197)	6091.6 (5328.7 to 6963.8)			
Opsono-14 (N=197)	4544.9 (4030.2 to 5125.4)			
Opsono-18C (N=196)	3827.5 (3367.5 to 4350.4)			
Opsono-19F (N=201)	1251 (1045.6 to 1496.8)			
Opsono-23F (N=198)	4629.1 (3890.6 to 5507.8)			

Statistical analyses

No statistical analyses for this end point

Secondary: Anti-6A and anti-19A antibody concentrations.

End point title	Anti-6A and anti-19A antibody concentrations. ^[19]
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End point description:

End point type	Secondary
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End point timeframe:

1 month after vaccination with the second dose of 10-valent pneumococcal conjugate vaccine (Month 12).

Notes:

[19] - The end point is not reporting statistics for all the arms in the baseline period. It is expected all the baseline period arms will be reported on when providing values for an end point on the baseline period.

Justification: This analysis was based on the unprimed group only.

End point values	Unprimed Group			
Subject group type	Reporting group			
Number of subjects analysed	209			
Units: µg/mL				
geometric mean (confidence interval 95%)				
Anti-6A	0.51 (0.43 to 0.6)			
Anti-19A	1.99 (1.68 to 2.36)			

Statistical analyses

No statistical analyses for this end point

Secondary: Opsono-6A and opsono-19A titres.

End point title	Opsono-6A and opsono-19A titres. ^[20]
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End point description:

End point type	Secondary
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End point timeframe:

1 month after vaccination with the second dose of 10-valent pneumococcal conjugate vaccine (Month 12).

Notes:

[20] - The end point is not reporting statistics for all the arms in the baseline period. It is expected all the baseline period arms will be reported on when providing values for an end point on the baseline period.

Justification: This analysis was based on the unprimed group only

End point values	Unprimed Group			
Subject group type	Reporting group			
Number of subjects analysed	196			
Units: Titres				
geometric mean (confidence interval 95%)				
Opsono-6A (N=193)	918.6 (742.7 to 1136.1)			
Opsono-19A (N=196)	597.6 (467.2 to 764.4)			

Statistical analyses

No statistical analyses for this end point

Secondary: Anti-PD antibody concentrations.

End point title	Anti-PD antibody concentrations. ^[21]
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End point description:

End point type	Secondary			
End point timeframe: 1 month after vaccination with the second dose of 10-valent pneumococcal conjugate vaccine (Month 12).				
Notes: [21] - The end point is not reporting statistics for all the arms in the baseline period. It is expected all the baseline period arms will be reported on when providing values for an end point on the baseline period. Justification: This analysis was based on the unprimed group only.				
End point values	Unprimed Group			
Subject group type	Reporting group			
Number of subjects analysed	209			
Units: EL.U/mL				
geometric mean (confidence interval 95%)				
Anti-PD	785.9 (695.7 to 887.7)			

Statistical analyses

No statistical analyses for this end point

Secondary: Rabbit complement-mediated Serum Bactericidal Activity assay against *Neisseria meningitidis* serogroup A (rSBAMenA), rSBA-MenC, rSBA-MenW-135 and rSBA-MenY titres.

End point title	Rabbit complement-mediated Serum Bactericidal Activity assay against <i>Neisseria meningitidis</i> serogroup A (rSBAMenA), rSBA-MenC, rSBA-MenW-135 and rSBA-MenY titres. ^[22]
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End point description:

End point type	Secondary			
End point timeframe: 25-36 months post-vaccination in previous study				
Notes: [22] - The end point is not reporting statistics for all the arms in the baseline period. It is expected all the baseline period arms will be reported on when providing values for an end point on the baseline period. Justification: This analysis was based on the unprimed group only.				
End point values	Unprimed Group			
Subject group type	Reporting group			
Number of subjects analysed	83			
Units: Titres				
geometric mean (confidence interval 95%)				
rSBA-MenA (N=66)	325.5 (243.9 to 434.5)			
rSBA-MenC (N=80)	63.6 (41.2 to 98)			
rSBA-MenY (N=83)	372.2 (270 to 513)			

rSBA-MenW-135 (N=83)	247.6 (190.7 to 321.5)			
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Statistical analyses

No statistical analyses for this end point

Secondary: Number of subjects reported with any and grade 3 solicited local symptoms.

End point title	Number of subjects reported with any and grade 3 solicited local symptoms. ^[23]
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End point description:

End point type	Secondary
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End point timeframe:

During the 4-day (Days 0-3) post-vaccination period.

Notes:

[23] - The end point is not reporting statistics for all the arms in the baseline period. It is expected all the baseline period arms will be reported on when providing values for an end point on the baseline period.

Justification: This analysis was based on the primed baseline groups.

End point values	10Pn-PD-DiT/Paracetamol Group	10Pn-PD-DiT Group		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	112	108		
Units: Subjects				
Any pain	75	67		
Grade 3 pain	7	5		
Any redness	58	60		
Grade 3 redness	13	12		
Any swelling	43	41		
Grade 3 swelling	11	11		

Statistical analyses

No statistical analyses for this end point

Secondary: Number of subjects reported with any and grade 3 solicited local symptoms.

End point title	Number of subjects reported with any and grade 3 solicited local symptoms. ^[24]
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End point description:

End point type	Secondary
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End point timeframe:

During the 4-day (Days 0-3) post-vaccination period.

Notes:

[24] - The end point is not reporting statistics for all the arms in the baseline period. It is expected all the baseline period arms will be reported on when providing values for an end point on the baseline period.

Justification: This analysis was based on the unprimed group only

End point values	Unprimed Group			
Subject group type	Reporting group			
Number of subjects analysed	223			
Units: Subjects				
Any pain, D1 (N=223)	150			
Grade 3 pain, D1 (N=223)	13			
Any redness, D1 (N=223)	102			
Grade 3 redness, D1 (N=223)	23			
Any swelling, D1 (N=223)	78			
Grade 3 swelling, D1 (N=223)	14			
Any pain, D2 (N=222)	121			
Grade 3 pain, D2 (N=222)	12			
Any redness, D2 (N=222)	93			
Grade 3 redness, D2 (N=222)	12			
Any swelling, D2 (N=222)	65			
Grade 3 swelling, D2 (N=222)	3			
Any pain, Across (N=223)	167			
Grade 3 pain, Across (N=223)	23			
Any redness, Across (N=223)	127			
Grade 3 redness, Across (N=223)	29			
Any swelling, Across (N=223)	101			
Grade 3 swelling, Across (N=223)	16			

Statistical analyses

No statistical analyses for this end point

Secondary: Number of subjects reported with any, grade 3 and related solicited general symptoms.

End point title	Number of subjects reported with any, grade 3 and related solicited general symptoms. ^[25]
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End point description:

End point type	Secondary
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End point timeframe:

During the 4-day (Days 0-3) post-vaccination period.

Notes:

[25] - The end point is not reporting statistics for all the arms in the baseline period. It is expected all the baseline period arms will be reported on when providing values for an end point on the baseline period.

Justification: This analysis was based on the primed baseline groups.

End point values	10Pn-PD-DiT/Paracetamol Group	10Pn-PD-DiT Group		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	112	108		
Units: Subjects				
Any drowsiness	41	34		
Related drowsiness	0	0		
Grade 3 drowsiness	26	26		
Any Irritability	33	26		
Related Irritability	1	0		
Grade 3 Irritability	21	16		
Any loss of appetite	23	15		
Related loss of appetite	1	1		
Grade 3 loss of appetite	13	10		
Any fever (Axillary/37.5°C)	12	8		
Grade 3 fever (Axillary/39.5°C)	0	1		
Related fever	10	5		

Statistical analyses

No statistical analyses for this end point

Secondary: Number of subjects reported with any, grade 3 and related solicited general symptoms.

End point title	Number of subjects reported with any, grade 3 and related solicited general symptoms. ^[26]
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End point description:

End point type	Secondary
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End point timeframe:

During the 4-day (Days 0-3) post-vaccination period.

Notes:

[26] - The end point is not reporting statistics for all the arms in the baseline period. It is expected all the baseline period arms will be reported on when providing values for an end point on the baseline period.

Justification: This analysis was based on the unprimed group only

End point values	Unprimed Group			
Subject group type	Reporting group			
Number of subjects analysed	223			
Units: Subjects				
Any drowsiness, D1 (N=223)	79			
Grade 3 drowsiness, D1 (N=223)	1			
Related drowsiness, D1 (N=223)	56			
Any Irritability, D1 (N=223)	69			
Grade 3 Irritability, D1 (N=223)	2			
Related irritability, D1 (N=223)	49			
Any loss of appetite, D1 (N=223)	47			
Grade 3 loss of appetite, D1 (N=223)	5			

Related loss of appetite, D1 (N=223)	32			
Any fever (Axillary/37.5°C), D1 (N=223)	23			
Grade 3 fever (Axillary/39.5°C), D1 (N=223)	1			
Related fever, D1 (N=223)	15			
Any drowsiness, D2 (N=222)	60			
Grade 3 drowsiness, D2 (N=222)	1			
Related drowsiness, D2 (N=222)	43			
Any Irritability, D2 (N=222)	59			
Grade 3 Irritability, D2 (N=222)	1			
Related irritability, D2 (N=222)	44			
Any loss of appetite, D2 (N=222)	26			
Grade 3 loss of appetite, D2 (N=222)	1			
Related loss of appetite, D2 (N=222)	16			
Any fever (Axillary/37.5°C), D2 (N=222)	9			
Grade 3 fever (Axillary/39.5°C), D2 (N=222)	1			
Related fever, D2 (N=222)	5			
Any drowsiness, Across (N=223)	102			
Grade 3 drowsiness, Across (N=223)	2			
Related drowsiness, Across (N=223)	73			
Any Irritability, Across (N=223)	98			
Grade 3 Irritability, Across (N=223)	3			
Related irritability, Across (N=223)	74			
Any loss of appetite, Across (N=223)	61			
Grade 3 loss of appetite, Across (N=223)	6			
Related loss of appetite, Across (N=223)	42			
Any fever (Axillary/37.5°C), Across (N=223)	30			
Grade 3 fever (Axillary/39.5°C), Across (N=223)	2			
Related fever, Across (N=223)	19			

Statistical analyses

No statistical analyses for this end point

Secondary: Number of subjects reported with unsolicited adverse events (AEs).

End point title	Number of subjects reported with unsolicited adverse events (AEs).
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End point description:

End point type	Secondary
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End point timeframe:

Within 31 days after each vaccination.

End point values	10Pn-PD-DiT/Paracetamol Group	10Pn-PD-DiT Group	Unprimed Group	
Subject group type	Reporting group	Reporting group	Reporting group	
Number of subjects analysed	112	108	223	
Units: Subjects				
Any AE(s)	24	29	73	

Statistical analyses

No statistical analyses for this end point

Secondary: Number of subjects reported with serious adverse events (SAEs).

End point title	Number of subjects reported with serious adverse events (SAEs).
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End point description:

End point type	Secondary
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End point timeframe:

During the entire study period.

End point values	10Pn-PD-DiT/Paracetamol Group	10Pn-PD-DiT Group	Unprimed Group	
Subject group type	Reporting group	Reporting group	Reporting group	
Number of subjects analysed	112	108	223	
Units: Subjects				
Any SAE(s)	0	0	0	

Statistical analyses

No statistical analyses for this end point

Secondary: Number of subjects with S.pneumoniae (vaccine serotypes) in nasopharyngeal swabs.

End point title	Number of subjects with S.pneumoniae (vaccine serotypes) in nasopharyngeal swabs. ^[27]
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End point description:

End point type	Secondary
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End point timeframe:

At 31-44 months of age and prior to the single/first dose of 10-valent pneumococcal conjugate vaccine and 40-48 months of age.

Notes:

[27] - The end point is not reporting statistics for all the arms in the baseline period. It is expected all the baseline period arms will be reported on when providing values for an end point on the baseline period.

Justification: This analysis was based on the pooled primed group that included the subjects in the remaining baseline groups.

End point values	Unprimed Group	Poled primed Group		
Subject group type	Reporting group	Subject analysis set		
Number of subjects analysed	210	216		
Units: Subjects				
31-44 months (N=216; 210)	40	21		
40-48 months (N=212; 208)	46	20		

Statistical analyses

No statistical analyses for this end point

Secondary: Number of subjects with S.pneumoniae (cross-reactive serotypes) in nasopharyngeal swabs.

End point title	Number of subjects with S.pneumoniae (cross-reactive serotypes) in nasopharyngeal swabs. ^[28]
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End point description:

End point type	Secondary
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End point timeframe:

At 31-44 months of age and prior to the single/first dose of 10-valent pneumococcal conjugate vaccine and 40-48 months of age.

Notes:

[28] - The end point is not reporting statistics for all the arms in the baseline period. It is expected all the baseline period arms will be reported on when providing values for an end point on the baseline period.

Justification: This analysis was based on the pooled primed group that included the subjects in the remaining baseline groups.

End point values	Unprimed Group	Poled primed Group		
Subject group type	Reporting group	Subject analysis set		
Number of subjects analysed	210	216		
Units: Subjects				
31-44 months (N=216; 210)	9	13		
40-48 months (N=212; 208)	19	18		

Statistical analyses

No statistical analyses for this end point

Secondary: Number of subjects with S.pneumoniae (non-vaccine and non-cross-reactive serotypes) in nasopharyngeal swabs.

End point title	Number of subjects with S.pneumoniae (non-vaccine and non-
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End point description:

End point type	Secondary
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End point timeframe:

At 31-44 months of age and prior to the single/first dose of 10-valent pneumococcal conjugate vaccine and 40-48 months of age.

Notes:

[29] - The end point is not reporting statistics for all the arms in the baseline period. It is expected all the baseline period arms will be reported on when providing values for an end point on the baseline period.

Justification: This analysis was based on the pooled primed group that included the subjects in the remaining baseline groups.

End point values	Unprimed Group	Poled primed Group		
Subject group type	Reporting group	Subject analysis set		
Number of subjects analysed	210	216		
Units: Subjects				
31-44 months (N=216; 210)	25	34		
40-48 months (N=212; 208)	27	35		

Statistical analyses

No statistical analyses for this end point

Secondary: Number of subjects with H. influenzae in nasopharyngeal swabs.

End point title	Number of subjects with H. influenzae in nasopharyngeal swabs. ^[30]
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End point description:

End point type	Secondary
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End point timeframe:

At 31-44 months of age and prior to the single/first dose of 10-valent pneumococcal conjugate vaccine and 40-48 months of age.

Notes:

[30] - The end point is not reporting statistics for all the arms in the baseline period. It is expected all the baseline period arms will be reported on when providing values for an end point on the baseline period.

Justification: This analysis was based on the pooled primed group that included the subjects in the remaining baseline groups.

End point values	Unprimed Group	Poled primed Group		
Subject group type	Reporting group	Subject analysis set		
Number of subjects analysed	210	216		
Units: Subjects				
31-44 months (N=216; 210)	43	54		
40-48 months (N=212; 208)	89	74		

Statistical analyses

No statistical analyses for this end point

Secondary: Number of subjects with S. pneumonia (vaccine serotypes) in nasopharyngeal swabs.

End point title	Number of subjects with S. pneumonia (vaccine serotypes) in nasopharyngeal swabs. ^[31]
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End point description:

End point type	Secondary
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End point timeframe:

At 31-44 months of age and prior to the single/first dose of 10-valent pneumococcal conjugate vaccine and 40-48 months of age.

Notes:

[31] - The end point is not reporting statistics for all the arms in the baseline period. It is expected all the baseline period arms will be reported on when providing values for an end point on the baseline period.

Justification: This analysis was based on the pooled primed group that included the subjects in the remaining baseline groups.

End point values	Unprimed Group	Poled primed Group		
Subject group type	Reporting group	Subject analysis set		
Number of subjects analysed	210	216		
Units: Subjects				
31-44 months (N=216; 210)	39	20		
40-48 months (N=212; 208)	41	18		

Statistical analyses

No statistical analyses for this end point

Secondary: Number of subjects with S. pneumonia (cross-reactive serotypes) in nasopharyngeal swabs.

End point title	Number of subjects with S. pneumonia (cross-reactive serotypes) in nasopharyngeal swabs. ^[32]
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End point description:

End point type	Secondary
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End point timeframe:

At 31-44 months of age and prior to the single/first dose of 10-valent pneumococcal conjugate vaccine and 40-48 months of age.

Notes:

[32] - The end point is not reporting statistics for all the arms in the baseline period. It is expected all the baseline period arms will be reported on when providing values for an end point on the baseline period.

Justification: This analysis was based on the pooled primed group that included the subjects in the remaining baseline groups.

End point values	Unprimed Group	Poled primed Group		
Subject group type	Reporting group	Subject analysis set		
Number of subjects analysed	210	216		
Units: Subjects				
31-44 months (N=216; 210)	9	13		
40-48 months (N=212; 208)	18	18		

Statistical analyses

No statistical analyses for this end point

Secondary: Number of subjects with S. pneumonia (non-vaccine and non-cross-reactive serotypes) in nasopharyngeal swabs.

End point title	Number of subjects with S. pneumonia (non-vaccine and non-cross-reactive serotypes) in nasopharyngeal swabs. ^[33]
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End point description:

End point type	Secondary
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End point timeframe:

At 31-44 months of age and prior to the single/first dose of 10-valent pneumococcal conjugate vaccine and 40-48 months of age.

Notes:

[33] - The end point is not reporting statistics for all the arms in the baseline period. It is expected all the baseline period arms will be reported on when providing values for an end point on the baseline period.

Justification: This analysis was based on the pooled primed group that included the subjects in the remaining baseline groups.

End point values	Unprimed Group	Poled primed Group		
Subject group type	Reporting group	Subject analysis set		
Number of subjects analysed	210	216		
Units: Subjects				
31-44 months (N=216; 210)	24	32		
40-48 months (N=212; 208)	25	33		

Statistical analyses

No statistical analyses for this end point

Secondary: Number of subjects with H. influenzae in nasopharyngeal swabs.

End point title	Number of subjects with H. influenzae in nasopharyngeal swabs. ^[34]
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End point description:

End point type	Secondary
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End point timeframe:

At 31-44 months of age and prior to the single/first dose of 10-valent pneumococcal conjugate vaccine and 40-48 months of age.

Notes:

[34] - The end point is not reporting statistics for all the arms in the baseline period. It is expected all the baseline period arms will be reported on when providing values for an end point on the baseline period.

Justification: This analysis was based on the pooled primed group that included the subjects in the remaining baseline groups.

End point values	Unprimed Group	Poled primed Group		
Subject group type	Reporting group	Subject analysis set		
Number of subjects analysed	210	216		
Units: Subjects				
31-44 months (N=216; 210)	35	49		
40-48 months (N=212; 208)	70	50		

Statistical analyses

No statistical analyses for this end point

Adverse events

Adverse events information

Timeframe for reporting adverse events:

Solicited symptoms: 4-day follow-up periods after vaccination (Days 0- 3), across doses; Unsolicited AEs: 31-day follow-up periods after vaccination (Days 0-30), across doses; SAEs: during the entire study period.

Assessment type	Non-systematic
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Dictionary used

Dictionary name	MedDRA
Dictionary version	14.0

Reporting groups

Reporting group title	10Pn-PD-DiT/ Paracetamol Group
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Reporting group description:

Subjects were vaccinated with three primary vaccination doses with prophylactic administration of paracetamol in study 10PN-PD-DIT-010 (107017), and a booster dose of GSK Biologicals' Synflorix™ vaccine, co-administered with Infanrix™ hexa and with prophylactic antipyretic treatment (rectal paracetamol) in study 10PN-PD-DIT-014 BST-010 and who received in this study one dose of Synflorix™ vaccine at month 9 (40-48 months of age).

Reporting group title	10Pn-PD-DiT Group
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Reporting group description:

Subjects were vaccinated with three primary vaccination doses with prophylactic administration of paracetamol in study 10PN-PD-DIT-010 (107017), at 12-15 months of age a booster dose of GSK Biologicals' Synflorix™ vaccine, co-administered with Infanrix™ hexa without prophylactic antipyretic treatment (rectal paracetamol) in study 10PN-PD-DIT-014 BST-010 (107137) and one dose of Synflorix™ vaccine in this study at Month 9 (month 40-48 of age).

Reporting group title	Unprimed Group
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Reporting group description:

Subjects were vaccinated with three primary vaccination doses without prophylactic administration of paracetamol in study 10PN-PD-DIT-010 (107017), at 12-15 months of age a booster dose of GSK Biologicals' 10Pn-PD-DiT vaccine, co-administered with Infanrix™ hexa without prophylactic antipyretic treatment (rectal) in study 10PN-PD-DIT-014 BST-010 (107137) and two doses, one of the 10Pn-PD-DiT vaccine at Month 9 (40-48 months of age) and one at Month 11 (42-50 months of age).

Serious adverse events	10Pn-PD-DiT/ Paracetamol Group	10Pn-PD-DiT Group	Unprimed Group
Total subjects affected by serious adverse events			
subjects affected / exposed	0 / 112 (0.00%)	0 / 108 (0.00%)	0 / 223 (0.00%)
number of deaths (all causes)	0	0	0
number of deaths resulting from adverse events	0	0	0

Frequency threshold for reporting non-serious adverse events: 5 %

Non-serious adverse events	10Pn-PD-DiT/ Paracetamol Group	10Pn-PD-DiT Group	Unprimed Group
Total subjects affected by non-serious adverse events			
subjects affected / exposed	75 / 112 (66.96%)	67 / 108 (62.04%)	167 / 223 (74.89%)
General disorders and administration site conditions			
Pain			
alternative assessment type: Systematic			
subjects affected / exposed	75 / 112 (66.96%)	67 / 108 (62.04%)	167 / 223 (74.89%)
occurrences (all)	75	67	167
Redness			
alternative assessment type: Systematic			
subjects affected / exposed	58 / 112 (51.79%)	60 / 108 (55.56%)	127 / 223 (56.95%)
occurrences (all)	58	60	127
Swelling			
alternative assessment type: Systematic			
subjects affected / exposed	43 / 112 (38.39%)	41 / 108 (37.96%)	101 / 223 (45.29%)
occurrences (all)	43	41	101
Drowsiness			
alternative assessment type: Systematic			
subjects affected / exposed	41 / 112 (36.61%)	34 / 108 (31.48%)	102 / 223 (45.74%)
occurrences (all)	41	34	102
Irritability			
alternative assessment type: Systematic			
subjects affected / exposed	33 / 112 (29.46%)	26 / 108 (24.07%)	98 / 223 (43.95%)
occurrences (all)	33	26	98
Loss of appetite			
alternative assessment type: Systematic			
subjects affected / exposed	23 / 112 (20.54%)	15 / 108 (13.89%)	61 / 223 (27.35%)
occurrences (all)	23	15	61
Fever/(Axillary $\geq 37.5^{\circ}\text{C}$)			
alternative assessment type: Systematic			
subjects affected / exposed	12 / 112 (10.71%)	8 / 108 (7.41%)	30 / 223 (13.45%)
occurrences (all)	12	8	30

More information

Substantial protocol amendments (globally)

Were there any global substantial amendments to the protocol? Yes

Date	Amendment
29 March 2011	<p>The primary objective of the current study is to demonstrate the immunological memory induced following primary vaccination in study 10PN-PD-DIT-010 (107017) and booster vaccination in study 10PN-PD-DIT-014 (107137) with the 10Pn-PD-DiT vaccine, through evaluation of early antibody responses after an additional dose of 10Pn-PD-DiT at 40-48 months of age. In addition, as secondary objective, the antibody persistence 25-36 months following vaccination with GSK Biologicals' MenACWYTT conjugate vaccine in study 10PN-PD-DIT-014 (107137) will be evaluated in terms of the percentage of subjects with <i>N. meningitidis</i> serogroup A, C, W-135 and Y titers ≥ 8 as measured by a serum bactericidal assay using rabbit complement (rSBA). In addition, to support the data obtained by rSBA testing, antibody concentrations against meningococcal polysaccharides are planned to be assessed by ELISA. However, the sponsor has decided not to perform the ELISA testing against meningococcal polysaccharides for this study for the following reasons:</p> <ul style="list-style-type: none">• the World Health Organization (WHO) considers SBA the primary means of assessing immune response to meningococcal conjugate vaccines [WHO, 2006; WHO, 1999]• circulating bactericidal antibodies are more critical for persistent protection against meningococcal disease than antibodies against meningococcal polysaccharides [CDC, 2011; WHO, 2006]. <p>Although antibody concentrations will not be determined by ELISA, all subjects will be informed of their antibody titers measured by rSBA when statistical analyses has been completed.</p>

Notes:

Interruptions (globally)

Were there any global interruptions to the trial? No

Limitations and caveats

None reported