



Clinical trial results:

An open-label study to assess the immune persistence in healthy Chinese toddlers primed in infancy with three doses of GSK Biologicals' DTPa-IPV/Hib vaccine, and to assess the safety and immunogenicity of a booster dose of IPV and DTPa/Hib administered at 18 to 24 months of age.

Summary

EudraCT number	2012-003324-20
Trial protocol	Outside EU/EEA
Global end of trial date	16 January 2012

Results information

Result version number	v1
This version publication date	01 April 2016
First version publication date	10 July 2015

Trial information

Trial identification

Sponsor protocol code	114386
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Additional study identifiers

ISRCTN number	-
ClinicalTrials.gov id (NCT number)	NCT01449812
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, 44 2989904466, GSKClinicalSupportHD@gsk.com
Scientific contact	Clinical Trials Call Center, GlaxoSmithKline Biologicals, 44 2989904466, GSKClinicalSupportHD@gsk.com

Notes:

Paediatric regulatory details

Is trial part of an agreed paediatric investigation plan (PIP)	No
Does article 45 of REGULATION (EC) No 1901/2006 apply to this trial?	No
Does article 46 of REGULATION (EC) No 1901/2006 apply to this trial?	Yes

Notes:

Results analysis stage

Analysis stage	Final
Date of interim/final analysis	16 January 2012
Is this the analysis of the primary completion data?	Yes
Primary completion date	16 January 2012
Global end of trial reached?	Yes
Global end of trial date	16 January 2012
Was the trial ended prematurely?	No

Notes:

General information about the trial

Main objective of the trial:

- To assess the persistence of antibodies to all vaccine antigens before the booster dose.
- To assess the immune response to the study vaccines in terms of seroprotection to diphtheria, tetanus, Haemophilus influenzae type b and poliovirus types 1, 2 and 3, and in terms of vaccine response to the pertussis antigens, one month after booster vaccination.
- To assess the immune response to the study vaccines in terms of antibody concentrations or titres for all antigens, one month after the booster dose.

Protection of trial subjects:

All subjects were supervised after vaccination/product administration with appropriate medical treatment readily available. Vaccines were administered by qualified and trained personnel. Vaccines were administered only to eligible subjects that had no contraindications to any components of the vaccines.

Background therapy: -

Evidence for comparator: -

Actual start date of recruitment	16 October 2011
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	China: 825
Worldwide total number of subjects	825
EEA total number of subjects	0

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)	825
Children (2-11 years)	0
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.

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	Infanrix+Hib/Poliorix 1 Group
Arm description: -	
Arm type	Experimental
Investigational medicinal product name	Infanrix+Hib™
Investigational medicinal product code	
Other name	
Pharmaceutical forms	Injection
Routes of administration	Intramuscular use

Dosage and administration details:

Subjects primed with 3 doses of the **Infanrix-IPV/Hib™** vaccine at 2, 3, 4 months of age in the primary 112584 study, received 1 dose of **Poliorix™** and of **Infanrix+Hib™** vaccines at 18-24 months of age. The **Poliorix™** and **Infanrix+Hib™** vaccines were administered as an intramuscular (IM) injection into the upper sides of the left and right thighs, respectively.

Investigational medicinal product name	Poliorix™
Investigational medicinal product code	
Other name	
Pharmaceutical forms	Injection
Routes of administration	Intramuscular use

Dosage and administration details:

Subjects primed with 3 doses of the **Infanrix-IPV/Hib™** vaccine at 2, 3, 4 months of age in the primary 112584 study, received 1 dose of **Poliorix™** and of **Infanrix+Hib™** vaccines at 18-24 months of age. The **Poliorix™** and **Infanrix+Hib™** vaccines were administered as an intramuscular (IM) injection into the upper sides of the left and right thighs, respectively.

Arm title	Infanrix+Hib/Poliorix 2 Group
Arm description: -	
Arm type	Experimental
Investigational medicinal product name	Infanrix+Hib™
Investigational medicinal product code	
Other name	
Pharmaceutical forms	Injection
Routes of administration	Intramuscular use

Dosage and administration details:

Subjects primed with 3 doses of the **Infanrix-IPV/Hib™** vaccine at 3, 4, 5 months of age in the primary 112584 study, received 1 dose of **Poliorix™** and of **Infanrix+Hib™** vaccines at 18-24 months of age. The **Poliorix™** and **Infanrix+Hib™** vaccines were administered as an intramuscular (IM) injection into the upper sides of the left and right thighs, respectively.

Investigational medicinal product name	Poliorix™
Investigational medicinal product code	
Other name	
Pharmaceutical forms	Injection
Routes of administration	Intramuscular use

Dosage and administration details:

Subjects primed with 3 doses of the Infanrix-IPV/Hib™ vaccine at 3, 4, 5 months of age in the primary 112584 study, received 1 dose of Poliorix™ and of Infanrix+Hib™ vaccines at 18-24 months of age. The Poliorix™ and Infanrix+Hib™ vaccines were administered as an intramuscular (IM) injection into the upper sides of the left and right thighs, respectively.

Arm title	Control Group
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Arm description: -

Arm type	Active comparator
Investigational medicinal product name	Infanrix+Hib™
Investigational medicinal product code	
Other name	
Pharmaceutical forms	Injection
Routes of administration	Intramuscular use

Dosage and administration details:

Subjects primed with 3 doses of the Infanrix+Hib™ vaccine at 2, 3, 4 months of age in the primary 112584 study, received 1 dose of Poliorix™ and of Infanrix+Hib™ vaccines at 18-24 months of age. The Poliorix™ and Infanrix+Hib™ vaccines were administered as an intramuscular (IM) injection into the upper sides of the left and right thighs, respectively.

Investigational medicinal product name	Poliorix™
Investigational medicinal product code	
Other name	
Pharmaceutical forms	Injection
Routes of administration	Intramuscular use

Dosage and administration details:

Subjects primed with 3 doses of the Infanrix+Hib™ vaccine at 2, 3, 4 months of age in the primary 112584 study, received 1 dose of Poliorix™ and of Infanrix+Hib™ vaccines at 18-24 months of age. The Poliorix™ and Infanrix+Hib™ vaccines were administered as an intramuscular (IM) injection into the upper sides of the left and right thighs, respectively.

Number of subjects in period 1	Infanrix+Hib/Poliorix 1 Group	Infanrix+Hib/Poliorix 2 Group	Control Group
Started	272	273	280
Completed	270	273	279
Not completed	2	0	1
Consent withdrawn by subject	-	-	1
Migrated/moved from study area	1	-	-
Lost to follow-up	1	-	-

Baseline characteristics

Reporting groups

Reporting group title	Infanrix+Hib/Poliorix 1 Group
Reporting group description: -	
Reporting group title	Infanrix+Hib/Poliorix 2 Group
Reporting group description: -	
Reporting group title	Control Group
Reporting group description: -	

Reporting group values	Infanrix+Hib/Poliorix 1 Group	Infanrix+Hib/Poliorix 2 Group	Control Group
Number of subjects	272	273	280
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	19.5	19.4	19.5
standard deviation	± 0.93	± 0.91	± 0.97
Gender categorical Units: Subjects			
Female	131	126	120
Male	141	147	160

Reporting group values	Total		
Number of subjects	825		
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	0 0 0 0 0 0 0 0		

Age continuous Units: months arithmetic mean standard deviation	-		
Gender categorical Units: Subjects			
Female	377		
Male	448		

End points

End points reporting groups

Reporting group title	Infanrix+Hib/Poliorix 1 Group
Reporting group description: -	
Reporting group title	Infanrix+Hib/Poliorix 2 Group
Reporting group description: -	
Reporting group title	Control Group
Reporting group description: -	

Primary: Number of seropositive subjects for anti-pertussis toxoid (anti-PT), anti-filamentous haemagglutinin (anti-FHA) and anti-diphtheria (anti-D).

End point title	Number of seropositive subjects for anti-pertussis toxoid (anti-PT), anti-filamentous haemagglutinin (anti-FHA) and anti-diphtheria (anti-D). ^[1]
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End point description:

End point type	Primary
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End point timeframe:

Before booster vaccination

Notes:

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

Justification: The analysis of the primary endpoint was descriptive i.e. no statistical hypothesis test was performed.

End point values	Infanrix+Hib/Poliorix 1 Group	Infanrix+Hib/Poliorix 2 Group	Control Group	
Subject group type	Reporting group	Reporting group	Reporting group	
Number of subjects analysed	0 ^[2]	0 ^[3]	0 ^[4]	
Units: subjects				

Notes:

[2] - The record will be updated when the results become available.

[3] - The record will be updated when the results become available.

[4] - The record will be updated when the results become available.

Statistical analyses

No statistical analyses for this end point

Secondary: Number of subjects with solicited local symptoms

End point title	Number of subjects with solicited local symptoms
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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

End point values	Infanrix+Hib/P oliorix 1 Group	Infanrix+Hib/P oliorix 2 Group	Control Group	
Subject group type	Reporting group	Reporting group	Reporting group	
Number of subjects analysed	270	273	279	
Units: subjects				
Any Pain	73	74	76	
Any Redness	19	15	19	
Any Swelling	16	10	14	

Statistical analyses

No statistical analyses for this end point

Secondary: Number of subjects with solicited general symptoms

End point title	Number of subjects with solicited general symptoms
End point description:	
End point type	Secondary
End point timeframe:	
During the 4-day (Days 0-3) post-vaccination period	

End point values	Infanrix+Hib/P oliorix 1 Group	Infanrix+Hib/P oliorix 2 Group	Control Group	
Subject group type	Reporting group	Reporting group	Reporting group	
Number of subjects analysed	270	273	279	
Units: subjects				
Any Drowsiness	38	50	38	
Any Irritability	78	81	72	
Any Loss of appetite	67	73	69	
Any Fever	102	105	91	

Statistical analyses

No statistical analyses for this end point

Secondary: Number of subjects with unsolicited AEs

End point title	Number of subjects with unsolicited AEs
End point description:	
End point type	Secondary

End point timeframe:

Within the 31-day (Days 0-30) period following booster vaccination

End point values	Infanrix+Hib/P oliorix 1 Group	Infanrix+Hib/P oliorix 2 Group	Control Group	
Subject group type	Reporting group	Reporting group	Reporting group	
Number of subjects analysed	272	273	280	
Units: subjects				
Any AEs	16	13	21	

Statistical analyses

No statistical analyses for this end point

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

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

End point type	Secondary
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End point timeframe:

Throughout the entire study period

End point values	Infanrix+Hib/P oliorix 1 Group	Infanrix+Hib/P oliorix 2 Group	Control Group	
Subject group type	Reporting group	Reporting group	Reporting group	
Number of subjects analysed	272	273	280	
Units: subjects				
Any SAEs	1	0	0	

Statistical analyses

No statistical analyses for this end point

Adverse events

Adverse events information

Timeframe for reporting adverse events:

Solicited local and general symptoms: during the 4-day (Days 0-3) post-vaccination period. AEs: within the 31-day (Days 0-30) period following booster vaccination. SAEs: throughout the entire study period.

Assessment type	Systematic
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Dictionary used

Dictionary name	MedDRA
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Dictionary version	14.1
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Reporting groups

Reporting group title	Infanrix+Hib/Poliorix 1 Group
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Reporting group description: -	
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Reporting group title	Infanrix+Hib/Poliorix 2 Group
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Reporting group description: -	
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Reporting group title	Control Group
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Reporting group description: -	
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Serious adverse events	Infanrix+Hib/Poliorix 1 Group	Infanrix+Hib/Poliorix 2 Group	Control Group
Total subjects affected by serious adverse events			
subjects affected / exposed	1 / 272 (0.37%)	0 / 273 (0.00%)	0 / 280 (0.00%)
number of deaths (all causes)	0	0	0
number of deaths resulting from adverse events			
Nervous system disorders			
Febrile convulsion			
subjects affected / exposed	1 / 272 (0.37%)	0 / 273 (0.00%)	0 / 280 (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
Infections and infestations			
Bronchopneumonia			
alternative assessment type: Non-systematic			
subjects affected / exposed	1 / 272 (0.37%)	0 / 273 (0.00%)	0 / 280 (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

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

Non-serious adverse events	Infanrix+Hib/Poliorix 1 Group	Infanrix+Hib/Poliorix 2 Group	Control Group
Total subjects affected by non-serious adverse events			
subjects affected / exposed	102 / 272 (37.50%)	105 / 273 (38.46%)	91 / 280 (32.50%)
General disorders and administration site conditions			
Pain			
subjects affected / exposed ^[1]	73 / 270 (27.04%)	74 / 273 (27.11%)	76 / 279 (27.24%)
occurrences (all)	73	74	76
Redness			
subjects affected / exposed ^[2]	19 / 270 (7.04%)	15 / 273 (5.49%)	16 / 279 (5.73%)
occurrences (all)	19	15	16
Swelling			
subjects affected / exposed ^[3]	16 / 270 (5.93%)	10 / 273 (3.66%)	14 / 279 (5.02%)
occurrences (all)	16	10	14
Drowsiness			
subjects affected / exposed ^[4]	38 / 270 (14.07%)	50 / 273 (18.32%)	38 / 279 (13.62%)
occurrences (all)	38	50	38
Irritability			
subjects affected / exposed ^[5]	78 / 270 (28.89%)	81 / 273 (29.67%)	72 / 279 (25.81%)
occurrences (all)	78	81	72
Loss of appetite			
subjects affected / exposed ^[6]	67 / 270 (24.81%)	73 / 273 (26.74%)	69 / 279 (24.73%)
occurrences (all)	67	73	69
Fever			
subjects affected / exposed ^[7]	102 / 270 (37.78%)	105 / 273 (38.46%)	91 / 279 (32.62%)
occurrences (all)	102	105	91
Infections and infestations			
Nasopharyngitis			
alternative assessment type: Non-systematic			
subjects affected / exposed	8 / 272 (2.94%)	6 / 273 (2.20%)	13 / 280 (4.64%)
occurrences (all)	8	6	13

Notes:

[1] - The number of subjects exposed to this adverse event is less than the total number of subjects exposed for the reporting group. These numbers are expected to be equal.

Justification: Solicited local and general symptoms were only reported for subjects with a symptom sheet completed.

[2] - The number of subjects exposed to this adverse event is less than the total number of subjects exposed for the reporting group. These numbers are expected to be equal.

Justification: Solicited local and general symptoms were only reported for subjects with a symptom sheet completed.

[3] - The number of subjects exposed to this adverse event is less than the total number of subjects exposed for the reporting group. These numbers are expected to be equal.

Justification: Solicited local and general symptoms were only reported for subjects with a symptom sheet completed.

[4] - The number of subjects exposed to this adverse event is less than the total number of subjects exposed for the reporting group. These numbers are expected to be equal.

Justification: Solicited local and general symptoms were only reported for subjects with a symptom sheet completed.

[5] - The number of subjects exposed to this adverse event is less than the total number of subjects exposed for the reporting group. These numbers are expected to be equal.

Justification: Solicited local and general symptoms were only reported for subjects with a symptom sheet completed.

[6] - The number of subjects exposed to this adverse event is less than the total number of subjects exposed for the reporting group. These numbers are expected to be equal.

Justification: Solicited local and general symptoms were only reported for subjects with a symptom sheet completed.

[7] - The number of subjects exposed to this adverse event is less than the total number of subjects exposed for the reporting group. These numbers are expected to be equal.

Justification: Solicited local and general symptoms were only reported for subjects with a symptom sheet completed.

More information

Substantial protocol amendments (globally)

Were there any global substantial amendments to the protocol? Yes

Date	Amendment
12 July 2011	Due to significant revisions to the Chinese Pharmacopeia, the DTPa-IPV/Hib vaccine can currently not be locally retested and released in that country. The study design is therefore being modified to boost all subjects with the DTPa/Hib (Infanrix Hib) and IPV (Poliorix) vaccines.

Notes:

Interruptions (globally)

Were there any global interruptions to the trial? No

Limitations and caveats

None reported