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## **Pneumococcal conjugate vaccines for preventing acute otitis media in children (Review)**

de Sévaux JLH, Venekamp RP, Lutje V, Hak E, Schilder AGM, Sanders EAM, Damoiseaux RAMJ

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## [Intervention Review]

# Pneumococcal conjugate vaccines for preventing acute otitis media in children

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## ABSTRACT

### Background

Prior to introducing pneumococcal conjugate vaccines (PCVs), *Streptococcus pneumoniae* was most commonly isolated from the middle ear fluid of children with acute otitis media (AOM). Reducing nasopharyngeal colonisation of this bacterium by PCVs may lead to a decline in AOM. The effects of PCVs deserve ongoing monitoring since studies from the post-PCV era report a shift in causative otopathogens towards non-vaccine serotypes and other bacteria. This updated Cochrane Review was first published in 2002 and updated in 2004, 2009, 2014, and 2019.

### Objectives

To assess the effect of PCVs in preventing AOM in children up to 12 years of age.

### Search methods

We searched CENTRAL, MEDLINE, Embase, CINAHL, LILACS, Web of Science, and two trials registers, ClinicalTrials.gov and WHO ICTRP, to 11 June 2020.

### Selection criteria

Randomised controlled trials of PCV versus placebo or control vaccine.

### Data collection and analysis

We used the standard methodological procedures expected by Cochrane. The primary outcomes were frequency of all-cause AOM and adverse effects. Secondary outcomes included frequency of pneumococcal AOM and frequency of recurrent AOM (defined as three or more AOM episodes in six months or four or more in one year). We used GRADE to assess the certainty of the evidence.

## Main results

We included 15 publications of 11 trials (60,733 children, range 74 to 37,868 per trial) of 7- to 11-valent PCVs versus control vaccines (meningococcus type C vaccine in three trials, and hepatitis A or B vaccine in eight trials). We included one additional publication of a previously included trial for this 2020 update. We did not find any relevant trials with the newer 13-valent PCV. Most studies were funded by pharmaceutical companies. Overall, risk of bias was low. In seven trials (59,415 children), PCVs were administered in early infancy, whilst four trials (1318 children) included children aged one year and over who were either healthy or had a history of respiratory illness. There was considerable clinical heterogeneity across studies, therefore we reported results from individual studies.

### PCV administered in early infancy

#### PCV7

The licenced 7-valent PCV with CRM197 as carrier protein (CRM197-PCV7) was associated with a 6% (95% confidence interval (CI) -4% to 16%; 1 trial; 1662 children) and 6% (95% CI 4% to 9%; 1 trial; 37,868 children) relative risk reduction (RRR) in low-risk infants (moderate-certainty evidence), but was not associated with a reduction in all-cause AOM in high-risk infants (RRR -5%, 95% CI -25% to 12%). PCV7 with the outer membrane protein complex of *Neisseria meningitidis* serogroup B as carrier protein (OMPC-PCV7) was not associated with a reduction in all-cause AOM (RRR -1%, 95% CI -12% to 10%; 1 trial; 1666 children; low-certainty evidence).

CRM197-PCV7 and OMPC-PCV7 were associated with 20% (95% CI 7% to 31%) and 25% (95% CI 11% to 37%) RRR in pneumococcal AOM, respectively (2 trials; 3328 children; high-certainty evidence), and CRM197-PCV7 with 9% (95% CI -12% to 27%) and 10% (95% CI 7% to 13%) RRR in recurrent AOM (2 trials; 39,530 children; moderate-certainty evidence).

#### PHiD-CV10/11

The effect of a licenced 10-valent PCV conjugated to protein D, a surface lipoprotein of *Haemophilus influenzae*, (PHiD-CV10) on all-cause AOM in healthy infants varied from 6% (95% CI -6% to 17%; 1 trial; 5095 children) to 15% (95% CI -1% to 28%; 1 trial; 7359 children) RRR (low-certainty evidence). PHiD-CV11 was associated with 34% (95% CI 21% to 44%) RRR in all-cause AOM (1 trial; 4968 children; moderate-certainty evidence).

PHiD-CV10 and PHiD-CV11 were associated with 53% (95% CI 16% to 74%) and 52% (95% CI 37% to 63%) RRR in pneumococcal AOM (2 trials; 12,327 children; high-certainty evidence), and PHiD-CV11 with 56% (95% CI -2% to 80%) RRR in recurrent AOM (1 trial; 4968 children; low-certainty evidence).

### PCV administered at a later age

#### PCV7

We found no evidence of a beneficial effect on all-cause AOM of administering CRM197-PCV7 in children aged 1 to 7 years with a history of respiratory illness or frequent AOM (2 trials; 457 children; moderate-certainty evidence) and CRM197-PCV7 combined with a trivalent influenza vaccine in children aged 18 to 72 months with a history of respiratory tract infections (1 trial; 597 children; moderate-certainty evidence).

#### CRM197-PCV9

In 1 trial including 264 healthy daycare attendees aged 1 to 3 years, CRM197-PCV9 was associated with 17% (95% CI -2% to 33%) RRR in parent-reported all-cause otitis media (very low-certainty evidence).

## Adverse events

Nine trials reported on adverse effects (77,389 children; high-certainty evidence). Mild local reactions and fever were common in both groups, and occurred more frequently in PCV than in control vaccine groups: redness (< 2.5 cm): 5% to 20% versus 0% to 16%; swelling (< 2.5 cm): 5% to 12% versus 0% to 8%; and fever (< 39 °C): 15% to 44% versus 8% to 25%. More severe redness (> 2.5 cm), swelling (> 2.5 cm), and fever (> 39 °C) occurred less frequently (0% to 0.9%, 0.1% to 1.3%, and 0.4% to 2.5%, respectively) in children receiving PCV, and did not differ significantly between PCV and control vaccine groups. Pain or tenderness, or both, was reported more frequently in PCV than in control vaccine groups: 3% to 38% versus 0% to 8%. Serious adverse events judged to be causally related to vaccination were rare and did not differ significantly between groups, and no fatal serious adverse event judged causally related to vaccination was reported.

## Authors' conclusions

Administration of the licenced CRM197-PCV7 and PHiD-CV10 during early infancy is associated with large relative risk reductions in pneumococcal AOM. However, the effects of these vaccines on all-cause AOM is far more uncertain based on low- to moderate-certainty evidence. We found no evidence of a beneficial effect on all-cause AOM of administering PCVs in high-risk infants, after early infancy, and in older children with a history of respiratory illness. Compared to control vaccines, PCVs were associated with an increase in mild local reactions (redness, swelling), fever, and pain and/or tenderness. There was no evidence of a difference in more severe local reactions, fever, or serious adverse events judged to be causally related to vaccination.

## PLAIN LANGUAGE SUMMARY

### Pneumococcal vaccination for preventing acute middle ear infections in children

#### Review question

We reviewed the evidence for the effect of vaccination against *Streptococcus pneumoniae* (pneumococcus, a type of bacterium) for preventing acute middle ear infections in children.

#### Background

Before nationwide implementation of vaccination against *S pneumoniae* with pneumococcal conjugate vaccines (PCVs), pneumococcus was the most frequent cause of acute middle ear infections in children. Vaccination against this bacterium with PCVs may therefore lead to fewer acute middle ear infections in children. However, ongoing monitoring of the effects of PCVs on acute middle ear infections is warranted, since recent studies report a shift in bacteria causing acute middle ear infections towards pneumococcal types not included in the vaccines and other bacteria.

#### Study characteristics

The evidence is current up to 11 June 2020. We included 11 trials of PCVs versus control vaccines (meningococcus type C conjugate vaccine in three trials, and hepatitis A or B vaccine in eight trials) involving a total of 60,733 children. The PCVs used in the trials contained 7 to 11 different types of pneumococcus. None of the trials used the newer PCV containing 13 different types. Most trials were funded by pharmaceutical companies. Overall, risk of bias was low. In seven trials (59,415 children), children received PCVs in early infancy, whilst four trials included 1318 children aged one year and over who were either healthy or who had previous respiratory illness.

#### Key results

When a licenced vaccine containing seven different types of pneumococcus (CRM197-PCV7) was given during early infancy, the risk of experiencing acute middle ear infections increased by 5% in high-risk infants and decreased by 6% in low-risk infants. When administering a licenced vaccine containing 10 types of pneumococcus together with a carrier protein from another bacterium called *Haemophilus influenzae* (PHiD-CV10), the risk of experiencing acute middle ear infections decreased by 6% to 15%, however neither of these estimates reached significance.

Giving PCV7 after early infancy (children aged one year and above) and in older children with a history of respiratory illness or frequent acute middle ear infections was not associated with reductions in acute middle ear infections.

Mild local reactions (redness, swelling), fever, and pain/tenderness were common and occurred more frequently in children receiving PCV than in those receiving control vaccines. More severe local reactions (redness and swelling > 2.5 cm) and fever (> 39 °C) occurred far less frequently and did not differ between vaccine groups. Serious adverse events judged to have been related to vaccination were rare and did not differ significantly between vaccine groups.

#### Certainty of the evidence

We assessed the certainty of the evidence for CRM197-PCV7 in early infancy to be moderate (further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate). We judged the certainty of the evidence for PHiD-CV10 to be low (further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate). We judged the certainty of the evidence for PCV7 in older children with or without a history of respiratory illness to be moderate (further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate).

## SUMMARY OF FINDINGS

### Summary of findings 1. Pneumococcal conjugate vaccine versus control vaccine for preventing acute otitis media in children

#### Pneumococcal conjugate vaccine versus control vaccine for preventing acute otitis media in children

**Patient or population:** infants (predominantly < 6 months of age) and older children (aged 1 to 7 years)

**Settings:** community (Finland, the Netherlands, the Czech Republic and Slovakia, Israel, the USA, Argentina, Colombia, and Panama)

**Intervention:** multivalent PCVs

**Comparison:** control vaccine

PCV type	VE - relative effect (95% CI)*	No. of participants (studies)	Certainty of evidence	Comments
<b>Frequency of all-cause acute otitis media</b>				
<b>CRM197-PCV7</b> in low-risk infants	RRR: 6% (−4% to 16%) to 6% (4% to 9%)#	39,530 (2 RCTs)	⊕⊕⊕⊕ Moderate <sup>a</sup>	Results are derived from 1 very large trial including 37,868 infants, <a href="#">Black 2000</a> /Fireman 2003, and 1 smaller trial including 1662 infants, <a href="#">Eskola 2001</a> /Palmu 2009.
<b>CRM197-PCV7</b> in high-risk infants	RRR: −5% (−25% to 12%)	944 (1 RCT)	⊕⊕⊕⊕ Low <sup>b</sup>	Results are derived from 1 relatively small trial with low risk of bias ( <a href="#">O'Brien 2008</a> ).
<b>OMPC-PCV7</b> in low-risk infants	RRR: −1% (−12% to 10%)	1666 (1 RCT)	⊕⊕⊕⊕ Low <sup>b</sup>	Results are derived from 1 trial with low risk of bias ( <a href="#">Kilpi 2003</a> ).
<b>PHiD-CV10</b> in low-risk infants	RRR: 6% (−6% to 17%) to 15% (−1% to 28%)	12,454 (2 RCTs)	⊕⊕⊕⊕ Low <sup>c</sup>	Results are derived from 2 trials with low, <a href="#">Tregnaghi 2014</a> /Sáez-Llorens 2017, and unclear risk of bias ( <a href="#">Vesikari 2016</a> /Karppinen 2019).  AOM incidence rate in the control group of 1 trial, <a href="#">Tregnaghi 2014</a> /Sáez-Llorens 2017, was low compared to other studies ( <a href="#">Table 1</a> ).
<b>PHiD-CV11</b> in low-risk infants	RRR: 34% (21% to 44%)	4968 (1 RCT)	⊕⊕⊕⊕ Moderate <sup>d</sup>	Results are derived from 1 trial with low risk of bias ( <a href="#">Prymula 2006</a> ).  AOM incidence rate in the control group was low compared to other studies ( <a href="#">Table 1</a> ).
<b>Adverse effects</b>				
<b>CRM197-PCV7</b> in low-risk infants	Mild local reactions and fever were common in both groups, occurring more frequently in the PCV than in the control vaccine groups: redness (< 2.5 cm): 5% to 20% versus 0% to 16%, swelling (< 2.5 cm): 5%	77,389 (9 RCTs)	⊕⊕⊕⊕ High	Results are derived from 9 trials with low risk of bias.
<b>OMPC-PCV7</b> in low-risk infants				

**PHiD-PC10 and PHiD-PC11 in low-risk infants**
**CRM197-PCV7/9 and CRM197-PCV7 plus TIV in older children**

to 12% versus 0% to 8%, and fever (< 39 °C): 15% to 44% versus 8% to 25%.

More severe redness (> 2.5 cm), swelling (> 2.5 cm), and fever (> 39 °C) occurred less frequently (0% to 0.9%, 0.1% to 1.3%, and 0.4% to 2.5%, respectively, in children receiving PCV) and did not differ significantly between PCV and control vaccine groups.

Pain/tenderness was reported more frequently in children receiving PCV than in those receiving control vaccines: 3% to 38% versus 0% to 8%.

Serious adverse events judged to be causally related to vaccination were rare and did not differ significantly between vaccine groups. No fatal serious adverse event judged to be causally related to vaccination was reported.

**Frequency of pneumococcal acute otitis media**

<b>CRM197-PCV7 in low-risk infants</b>	RRR: 20% (7% to 31) to 34% (21% to 45%)	1662 (1 RCT)	⊕⊕⊕⊕ High	Results are derived from 1 trial with low risk of bias ( <a href="#">Eskola 2001</a> /Palmu 2009).
<b>OMPC-PCV7 in low-risk infants</b>	RRR: 25% (11% to 37%)	1666 (1 RCT)	⊕⊕⊕⊕ High	Results are derived from 1 trial with low risk of bias ( <a href="#">Kilpi 2003</a> ).
<b>PHiD-CV10 in low-risk infants</b>	RRR: 53% (16% to 74%)	7359 (1 RCT)	⊕⊕⊕⊕ High	Results are derived from 1 trial with low risk of bias ( <a href="#">Tregnaighi 2014</a> /Sáez-Llorens 2017).
<b>PHiD-CV11 in low-risk infants</b>	RRR: 52% (37% to 63%)	4968 (1 RCT)	⊕⊕⊕⊕ High	Results are derived from 1 trial with low risk of bias ( <a href="#">Prymula 2006</a> ).

**Frequency of recurrent acute otitis media (defined as 3 or more acute otitis media episodes in 6 months or 4 or more in 1 year)**

<b>CRM197-PCV7 in low-risk infants</b>	RRR: 9% (-12% to 27%) to 10% (7% to 13%)	39,530 (2 RCTs)	⊕⊕⊕⊕ Moderate <sup>e</sup>	Results are derived from 1 very large trial including 37,868 infants, <a href="#">Black 2000</a> /Fireman 2003, and 1 smaller trial including 1662 infants, <a href="#">Eskola 2001</a> /Palmu 2009, both with low risk of bias.
<b>PHiD-CV11 in low-risk infants</b>	RRR: 56% (-2% to 80%)	4968 (1 RCT)	⊕⊕⊕⊕ Low <sup>f</sup>	Results are derived from 1 trial with low risk of bias ( <a href="#">Prymula 2006</a> ).

\*For readability purposes, absolute rates (episodes/person-year and incidence rate differences) are displayed in [Table 1](#).

#Depending on whether the outcome was assessed by a composite of positive culture and positive pneumolysin polymerase chain reaction (PCR) or by positive culture only, or whether ITT or per-protocol analysis was performed.

**GRADE (certainty in the evidence)**

**High certainty:** Further research is very unlikely to change our confidence in the estimate of effect.

**Moderate certainty:** Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

**Low certainty:** Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

**Very low certainty:** Any estimate of effect is very uncertain.

AOM: acute otitis media

CI: confidence interval

CRM197-PCV7: 7-valent pneumococcal conjugate vaccine conjugated to carrier protein CRM197

OMPC-PCV7: 7-valent pneumococcal conjugate vaccine conjugated to the outer membrane protein complex of *Neisseria meningitidis* serogroup B

PCV: pneumococcal conjugate vaccine

PHiD-CV10: 10-valent pneumococcal conjugate vaccine conjugated to protein D (surface lipoprotein of non-typeable *Haemophilus influenzae*)

PHiD-CV11: 11-valent pneumococcal conjugate vaccine conjugated to protein D (surface lipoprotein of non-typeable *Haemophilus influenzae*)

RCT: randomised controlled trial

RRR: relative risk reduction

TIV: trivalent influenza vaccine

VE: vaccine efficacy

<sup>a</sup>We downgraded the certainty of the evidence from high to moderate due to imprecise effect estimate and study limitations (risk of bias).

<sup>b</sup>We downgraded the certainty of the evidence from high to low due to the very imprecise effect estimate.

<sup>c</sup>We downgraded the certainty of the evidence from high to low due to study limitations (risk of bias) and imprecise effect estimates.

<sup>d</sup>We downgraded the certainty of the evidence from high to moderate due to indirectness of evidence (low AOM incidence rate in the control group compared to other studies, most likely due to methodological differences with other studies).

<sup>e</sup>We downgraded the certainty of the evidence from high to moderate due to imprecise effect estimate.

<sup>f</sup>We downgraded the certainty of the evidence from high to low due to imprecise effect estimate and indirectness of evidence (low AOM incidence rate in the control group compared to other studies, most likely due to methodological differences with other studies).



## BACKGROUND

### Description of the condition

Acute otitis media (AOM), defined as the presence of middle ear fluid together with one or more signs or symptoms of acute middle ear inflammation such as otalgia, otorrhoea, fever, or irritability, is one of the most common diseases in childhood and imposes a large burden on public health (Lieberthal 2013). Global AOM incidence rates are highest in children 1 to 4 years of age, with a peak incidence in 6- to 11-month-old infants (Monasta 2012). By the age of two years, up to 5% of all children have experienced recurrent AOM, defined as three or more AOM episodes in six months, or four or more in one year (Kvaerner 1997; Lieberthal 2013). The three main bacterial pathogens isolated from the middle ear fluid of children with AOM collected before the widespread use of pneumococcal conjugate vaccines (PCVs) were *Streptococcus pneumoniae* (25% to 39%), (non-typeable) *Haemophilus influenzae* (12% to 23%), and *Moraxella catarrhalis* (4% to 15%) (Bluestone 1992; Heikkinen 1999; Jacobs 1998; Luotonen 1981). Recent studies have shown that nationwide implementation of PCVs may have changed the frequency of the causative otopathogens involved in AOM towards pneumococcal serotypes not included in the vaccines and other bacteria including non-typeable *H influenzae* (Allemann 2017; Barenkamp 2017; Ben-Shimol 2019; Casey 2013; Coker 2010; Kaur 2017; Somech 2011; Tamir 2015; Wiertsema 2011).

### Description of the intervention

The marginal benefits of antibiotics for AOM in low-risk populations (Rovers 2006; Venekamp 2015); the increasing problem of bacterial resistance against antibiotics (Laxminarayan 2013); and the high estimated direct and indirect annual costs associated with AOM have prompted a search for effective vaccines to prevent this condition (Ahmed 2014; Boonacker 2011). With *S pneumoniae* (pneumococcus) being a common causative pathogen in childhood AOM and pneumonia, and one of the most frequent causes of invasive bacterial disease such as bacteraemia and meningitis, research has focused on the prevention of pneumococcal infections by pneumococcal vaccines. Pneumococcal polysaccharide vaccines (PPVs) have been available for decades, but have been shown to be poorly immunogenic in children aged up to two years, who are most prone to pneumococcal infections. In the most recent versions of this review, no further attention has been paid to the effect of PPVs, which were described in prior versions of this review (Straetemans 2003).

The first pneumococcal conjugate vaccines (PCVs), in which the pneumococcal capsular serotypes are covalently conjugated to carrier proteins, were developed in the 1990s and proved to be adequately immunogenic in infants and toddlers (Dagan 1997; Eskola 1999; Shinefield 1999). Over the past decades, various PCVs have been developed for use in children including:

- licenced 7-valent PCV containing the polysaccharides of seven serotypes (4, 6B, 9V, 14, 18C, 19F, and 23F) conjugated to the diphtheria-derived carrier protein CRM197 (**CRM197-PCV7**);
- 7-valent PCV with the outer membrane complex of *Neisseria meningitidis* serogroup B as carrier protein (**OMPC-PCV7**);
- 9-valent PCV containing the capsular polysaccharides of serotypes 1 and 5 in addition to those included in PCV7, conjugated to CRM197 (**CRM197-PCV9**);

- licenced 10-valent PCV containing the capsular polysaccharides of 10 serotypes (1, 4, 5, 6B, 7F, 9V, 14, 18C, 19F, and 23F) mostly conjugated to protein D, which is a surface lipoprotein of *H influenzae* (**PHiD-CV10**);
- 11-valent containing the capsular polysaccharides of serotype 3 as well as those included in PHiD-CV10 (**PHiD-CV11**); and
- licenced 13-valent PCV containing the capsular polysaccharides of 13 serotypes (1, 3, 4, 5, 6A, 6B, 7F, 9V, 14, 18C, 19A, 19F, and 23F) conjugated to CRM197 (**CRM197-PCV13**).

### How the intervention might work

Early and dense colonisation of the nasopharynx with bacterial otopathogens, including *S pneumoniae*, increases the risk of AOM substantially (Faden 1997; Leach 1994; Schilder 2016). As a consequence, reducing or eliminating nasopharyngeal colonisation of *S pneumoniae* by PCVs may lead to reductions in AOM incidence. In recent years, evidence has accumulated that PCVs might also disrupt the continuum of evolution from pneumococcal-associated otitis media (OM) towards chronic/recurrent OM by prevention of early vaccine-serotype AOM and thereby reducing subsequent and more complex disease caused by non-vaccine serotypes and non-typeable *H influenzae* (Ben-Shimol 2014; Dagan 2016).

### Why it is important to do this review

With AOM amongst the most common diseases in early childhood, the need for a vaccine to effectively prevent AOM is high. Over the past decades various randomised controlled trials have been performed to assess the effects of pneumococcal vaccination to prevent AOM. From 2009 onwards, two multivalent PCVs (PHiD-CV10 and CRM197-PCV13) have been licenced and are being implemented in nationwide immunisation programmes worldwide (WHO 2012). These new vaccines may have an increased benefit in preventing AOM (Marom 2014; O'Brien 2009; Soysal 2020). As such, it was important to provide an up-to-date systematic review on the effects of PCVs on preventing AOM. This review is an update of a Cochrane Review first published in 2002 (Straetemans 2002), and updated in 2004 (Straetemans 2004), 2009 (Jansen 2009), 2014 (Fortanier 2014), and 2019 (Fortanier 2019).

## OBJECTIVES

To assess the effect of PCVs in preventing AOM in children up to 12 years of age.

## METHODS

### Criteria for considering studies for this review

#### Types of studies

Randomised controlled trials (RCTs), irrespective of type, assessing the effect of pneumococcal conjugate vaccines (PCV) versus placebo or control vaccine in preventing acute otitis media (AOM) with a minimum follow-up duration of six months. As per previous versions of this review, we excluded studies that did not report outcome data relevant to this review.

#### Types of participants

Children aged up to 12 years.

## Types of interventions

PCV versus placebo or control vaccine.

## Types of outcome measures

We extracted data for the following predefined outcomes of interest.

### Primary outcomes

1. Frequency of all-cause AOM episodes, defined as AOM irrespective of causative pathogen. We considered this to be the most relevant outcome for children, parents, and clinicians.
2. Adverse effects including local (redness, swelling) and systemic reactions (fever), pain/tenderness, and serious adverse events (SAEs) judged to be causally related to vaccination.

### Secondary outcomes

1. Frequency of pneumococcal AOM.
2. Frequency of pneumococcal serotype-specific AOM (including vaccine serotype, non-vaccine serotype, and cross-reactive serotypes which are non-vaccine serotypes with a serogroup that is included in the vaccine).
3. Frequency of recurrent AOM (defined as three or more episodes in six months or four or more in one year).

## Search methods for identification of studies

The Cochrane Acute Respiratory Infections Group (2018 search update) and Cochrane Infectious Disease Group (2019 and 2020 search update) Information Specialists conducted systematic searches for RCTs and controlled clinical trials. There were no language, publication year, or publication status restrictions. The latest search was conducted on 11 June 2020.

### Electronic searches

For the 2014 review update, we used the search strategy presented in [Appendix 1](#).

For the 2020 and 2019 updates, we searched the Cochrane Central Register of Controlled Trials (CENTRAL; Issue 6, 2020), which contains the Cochrane Acute Respiratory Infections Specialised Register; MEDLINE (Ovid) (1995 to 11 June 2020); Embase (Elsevier) (1995 to 11 June 2020); CINAHL (EBSCO) (Cumulative Index to Nursing and Allied Health Literature) (2007 to 11 June 2020); LILACS (BIREME) (Latin American and Caribbean Health Science Information database) (2007 to 11 June 2020), and Science Citation Index Expanded (SCI-EXPANDED), Conference Proceedings Citation Index-Science (CPCI-S), and Current Chemical Reactions (CCR-EXPANDED) (all three from the Web of Science; Clarivate Analytics) (2007 to 11 June 2020).

We used the search strategy presented in [Appendix 2](#) to search CENTRAL and MEDLINE. We combined the MEDLINE search with the Cochrane Highly Sensitive Search Strategy for identifying randomised trials in MEDLINE: sensitivity- and precision-maximising version (2008 revision); Ovid format ([Lefebvre 2011](#)). We adapted the search strategy to search Embase ([Appendix 3](#)), CINAHL ([Appendix 4](#)), LILACS ([Appendix 5](#)), and Web of Science ([Appendix 6](#)).

## Searching other resources

To increase the yield of relevant studies, two review authors (JLHdS, RPV for the 2020 update) reviewed the reference lists of all relevant studies and review articles retrieved. We searched the US National Institutes of Health Ongoing Trials Register [ClinicalTrials.gov](https://clinicaltrials.gov/) ([clinicaltrials.gov/](#)) ([Appendix 7](#)) and the World Health Organization International Clinical Trials Registry Platform (WHO ICTRP) ([www.who.int/trialsearch](http://www.who.int/trialsearch)) ([Appendix 8](#)) on 4 September 2020 for completed and ongoing trials. We also searched the internet (via Google using the search terms 'pneumococcal conjugate vaccination for acute otitis media trial') and the extended abstracts published in the Recent Advances in Otitis Media (grey literature) on 4 September 2020 for any additional trials.

## Data collection and analysis

### Selection of studies

Two review authors (JLHdS, RPV for the 2020 update) independently screened the titles and abstracts identified by the database searches and reviewed the full text of those titles and abstracts deemed potentially relevant against the inclusion criteria. Any disagreements were resolved by discussion.

### Data extraction and management

Two review authors (JLHdS, RPV for the 2020 update) independently extracted data from the included studies. Any disagreements were resolved by discussion.

### Assessment of risk of bias in included studies

Two review authors (JLHdS, RPV for the 2020 update) independently assessed the methodological quality of the included trials. Any disagreements were resolved by discussion. We assessed the methodological quality of included studies using the 'Risk of bias' tool as described in Chapter 8 of the *Cochrane Handbook for Systematic Reviews of Interventions* ([Higgins 2011](#)). We judged the following domains as low, high, or unclear risk of bias: random sequence generation (selection bias), concealment of allocation (selection bias), blinding of participants and personnel (performance bias), blinding of outcome assessment (detection bias), incomplete outcome data (attrition bias), selective reporting (reporting bias), and other sources of bias.

### Measures of treatment effect

We expressed estimates of treatment effects as relative risks/hazard ratios with accompanying 95% confidence intervals (CIs). Vaccine efficacy was estimated as 1 minus the relative risk/hazard ratio (relative risk reduction (RRR)).

### Unit of analysis issues

We included all types of RCTs. In the case of cluster-randomised trials, we considered potential differences between the intervention effects being estimated and checked whether clustering was taken into account in the analysis of the individual trials.

### Dealing with missing data

For each trial, we determined the number of missing data and whether the authors took duration of follow-up (and censoring) of individual participants into account in their statistical analyses.

## Assessment of heterogeneity

We first assessed clinical heterogeneity across trials by reviewing the differences in the types of participants recruited, interventions used, and outcomes measured. We did not pool studies where clinical heterogeneity made it of no use to do so. Where studies were sufficiently homogeneous, we proposed to assess statistical heterogeneity for each outcome by visually inspecting the forest plots and by using the  $\chi^2$  test and the  $I^2$  statistic.

## Assessment of reporting biases

We proposed to assess reporting bias as within-study (outcome reporting) and between-study reporting (publication) bias (Higgins 2011).

### Outcome reporting bias

We searched the internet, ClinicalTrials.gov, and the WHO ICTRP for available study protocols to determine whether the outcomes reported in the studies were predefined, and whether all outcomes listed in the study protocol were reported in the trial publications. Where information was insufficient to judge risk of bias, we classified the risk of bias as unclear (Higgins 2011).

### Publication bias

We proposed a more formal method of assessing reporting bias, that is by creating funnel plots, if sufficient trials (10 or more) were available for a given outcome.

## Data synthesis

We primarily analysed the available data according to the intention-to-treat principle, that is by analysing all participants in the groups to which they were originally randomised. As a secondary analysis, we presented data based on a per-protocol analysis.

Where possible, we proposed conducting meta-analyses using Review Manager 5 by calculating treatment effects with the Mantel-Haenszel method, using a fixed-effect model where no substantial statistical heterogeneity was present ( $I^2 < 50\%$ ) (Review Manager 2014). If substantial statistical heterogeneity was detected and unresolved by sensitivity analysis, we proposed to calculate treatment effects using a random-effects (DerSimonian and Laird) model to provide more conservative effect estimates. Where clinical heterogeneity precluded meta-analyses, we reported the effect estimates as presented by the individual trials. If possible, we reported the incidences of the various outcomes in the study arms together with the vaccine efficacy estimates, with 95% CIs.

We proposed the following methods to conduct meta-analyses. The generalised Cox proportional hazard method proposed by Andersen 1982 is regarded as the most appropriate to assess the effect of PCVs on AOM (Jahn-Eimermacher 2007). Under the assumption that the hazard rate is proportional between both groups over time, and that the risk of AOM is not affected by previous episodes (although this is untrue), this model takes all available information into account, that is all episodes (including recurrences), differences in individual patient follow-up time, and time until a case of AOM (Jahn-Eimermacher 2007). However, information on individual follow-up time until the first, second, third, etc. case of AOM is difficult to obtain for each study to be included in the meta-analysis. Poisson regression is based on the assumption of a constant risk of AOM over time, and that this risk

is not affected by previous episodes of AOM. This method only requires the total follow-up time and total number of episodes, and therefore appears to be a more feasible method for meta-analysis. Furthermore, Poisson regression seems not to be affected by the deviation from a constant risk over time, having very similar results for the effect of PCVs on AOM to the Andersen-Gill approach (Jahn-Eimermacher 2007). For Poisson regression, the treatment effect is measured as a rate ratio defined as follows: (total AOM episodes in pneumococcal vaccination group divided by the number of children in the pneumococcal vaccination group multiplied by the follow-up time in months) divided by (total AOM episodes in control group divided by the number of children in the control group multiplied by the follow-up time in months) (McCullagh 1989).

## Subgroup analysis and investigation of heterogeneity

Because the effect of PCVs on AOM may be influenced by the age at which the PCV was administered, occurrence of previous AOM or respiratory tract infection episodes, and by the type of PCV used, we described the studies accordingly, that is we stratified those with vaccination in early infancy versus those with vaccination later in childhood by type of PCV used.

## Sensitivity analysis

We planned to carry out sensitivity analyses for risk of bias of the included studies to assess the robustness of review findings by excluding studies with high risk of bias (defined as high risk of bias for allocation concealment and high risk of attrition bias (overall loss to follow-up of more than 20% or differential follow-up observed, or both)) from meta-analysis.

## Summary of findings and assessment of the certainty of the evidence

We created [Summary of findings 1](#) for PCVs administered in early infancy using the following outcomes: frequency of all-cause AOM episodes, adverse effects, frequency of pneumococcal AOM, and frequency of recurrent AOM (defined as three or more AOM episodes in six months or four or more in one year). We used the five GRADE considerations (study limitations, consistency of effect, imprecision, indirectness, and publication bias) to assess the certainty of a body of evidence as it relates to the studies that contribute data to the meta-analyses for the prespecified outcomes (Atkins 2004). We judged the certainty of the evidence as high, moderate, low, or very low. We assessed evidence from RCTs that did not have serious limitations as of high certainty. However, we downgraded the certainty of evidence to moderate, low, or very low based on the following factors: study limitations (risk of bias), inconsistency (consistency of results), imprecision (precision of results), indirectness of evidence (directness of evidence), and publication bias (existence of publication bias). We used the methods and recommendations described in Chapter 14 of the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2019), employing GRADEpro GDT software (GRADEpro GDT). We justified all decisions to down- or upgrade the certainty of evidence using footnotes, and made comments to aid the reader's understanding of the review where necessary.

## RESULTS

### Description of studies

See [Characteristics of included studies](#) and [Characteristics of excluded studies](#).

### Results of the search

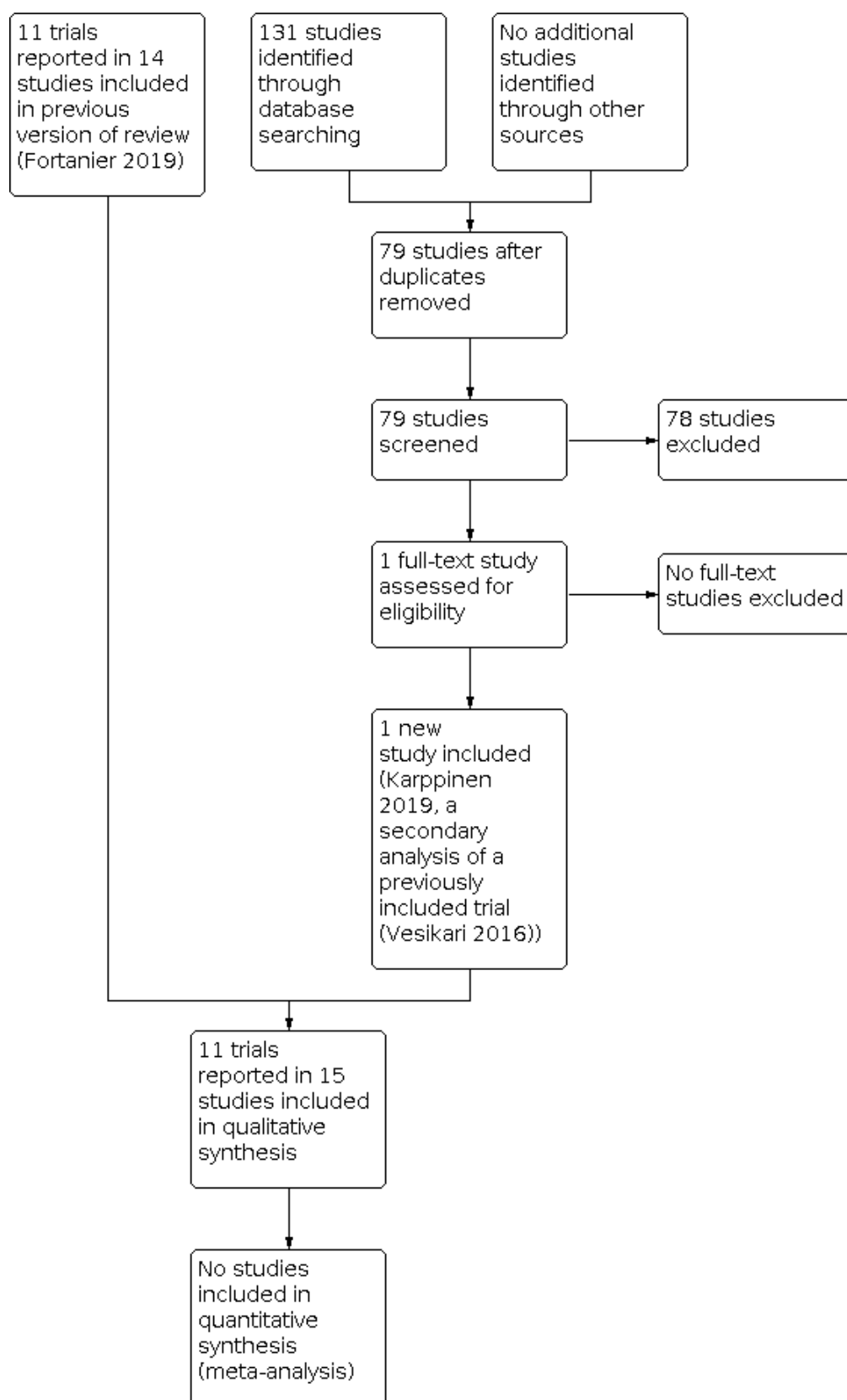
This review is an update of a Cochrane Review first published in 2002 ([Straetemans 2002](#)), and updated in 2004 ([Straetemans 2004](#)), 2009 ([Jansen 2009](#)), 2014 ([Fortanier 2014](#)), and 2019 ([Fortanier 2019](#)). In the 2019 review, which included studies up to March 2019, we included 11 RCTs reported in 14 publications ([Black 2000](#)/Fireman 2003; [Dagan 2001](#); [Eskola 2001](#)/Palmu 2009; [Jansen 2008](#); [Kilpi 2003](#); [O'Brien 2008](#); [Prymula 2006](#); [Tregnaghi 2014](#)/Sáez-Llorens 2017; [van Kempen 2006](#); [Veenhoven 2003](#); [Vesikari 2016](#)).

For the 2019 update, we searched electronic databases (December 2013 to March 2019) and retrieved 374 records. After removal of duplicates, we assessed 225 records by title and abstract, and

identified eight potentially eligible studies, which we obtained in full text. After reviewing the full texts, we excluded two publications that were additional analyses of the [Eskola 2001](#) study but did not include new outcome data useful to this review ([Palmu 2015a](#); [Sarasoja 2013](#)), and three publications that were secondary analyses of the Finnish invasive pneumococcal disease (FinIP) vaccine trial but did not report on any of our outcomes of interest ([Palmu 2014](#); [Palmu 2015b](#); [Palmu 2018](#)). This left three publications, Sáez-Llorens 2017; [Tregnaghi 2014](#); [Vesikari 2016](#), relating to two RCTs, [Tregnaghi 2014](#); [Vesikari 2016](#), suitable for inclusion. Sáez-Llorens 2017 was a further analysis of [Tregnaghi 2014](#).

For this 2020 update, we searched electronic databases (March 2019 to June 2020) and retrieved 131 records. After de-duplication, we assessed 79 unique records by title and abstract, and identified one additional potentially eligible study, which we obtained in full text. This study ([Karppinen 2019](#)) was suitable for inclusion and was included as secondary analysis of [Vesikari 2016](#) (FinIP vaccine trial). See [Figure 1](#).

**Figure 1. Study flow diagram.**





We did not identify additional relevant completed trials or any ongoing studies by scanning the reference lists of relevant systematic reviews or by searching the internet, the grey literature, or ClinicalTrials.gov and WHO ICTRP.

## Included studies

See [Characteristics of included studies](#) table.

We included 11 RCTs reported in 15 publications ([Black 2000](#)/[Fireman 2003](#); [Dagan 2001](#); [Eskola 2001](#)/[Palmu 2009](#); [Jansen 2008](#); [Kilpi 2003](#); [O'Brien 2008](#); [Prymula 2006](#); [Tregnaghi 2014](#)/[Sáez-Llorens 2017](#); [van Kempen 2006](#); [Veenhoven 2003](#); [Vesikari 2016](#)/[Karppinen 2019](#)). We added two RCTs (reported in four publications) for the 2019 and 2020 updates ([Tregnaghi 2014](#)/[Sáez-Llorens 2017](#); [Vesikari 2016](#)/[Karppinen 2019](#)). The included trials involved a total of 60,733 children.

## Study designs

Of the 11 included studies, nine were standard, individually randomised trials, and two were cluster-RCTs ([O'Brien 2008](#); [Vesikari 2016](#)). Both cluster-RCTs took the cluster effect into account in their analyses.

## Study populations (early infancy versus later in life)

In seven trials ([Black 2000](#)/[Fireman 2003](#); [Eskola 2001](#)/[Palmu 2009](#); [Kilpi 2003](#); [O'Brien 2008](#); [Prymula 2006](#); [Tregnaghi 2014](#)/[Sáez-Llorens 2017](#); [Vesikari 2016](#)/[Karppinen 2019](#)), PCVs were predominantly administered in children's first six months of life. Four trials assessed the effects of PCVs in children aged one year and over who were either healthy ([Dagan 2001](#)), or had a history of respiratory illness ([Jansen 2008](#); [van Kempen 2006](#); [Veenhoven 2003](#)). Three trials were conducted in Finland ([Eskola 2001](#)/[Palmu 2009](#); [Kilpi 2003](#); [Vesikari 2016](#)/[Karppinen 2019](#)), two in the USA ([Black 2000](#)/[Fireman 2003](#); [O'Brien 2008](#)), two in the Netherlands ([Jansen 2008](#); [Veenhoven 2003](#)), and the remaining in Belgium ([van Kempen 2006](#)), Israel ([Dagan 2001](#)), the Czech Republic and Slovakia ([Prymula 2006](#)), and Argentina, Colombia, and Panama ([Tregnaghi 2014](#)/[Sáez-Llorens 2017](#)). Most of these countries had AOM diagnosis and management guidelines at the time of the study ([Tamir 2017](#)).

## Interventions

### Type of PCV used and co-administration of other vaccines

In six trials, CRM197-PCV7 was used as the intervention ([Black 2000](#)/[Fireman 2003](#); [Eskola 2001](#)/[Palmu 2009](#); [Jansen 2008](#); [O'Brien 2008](#); [van Kempen 2006](#); [Veenhoven 2003](#)). In two studies, a booster dose with 23-valent PPV (containing capsular polysaccharides of the serotypes 1, 2, 3, 4, 5, 6B, 7F, 8, 9N, 9V, 10A, 11A, 12F, 14, 15B, 17F, 18C, 19F, 19A, 20, 22F, 23F, and 33F) was given to all children ([van Kempen 2006](#); [Veenhoven 2003](#)). In one trial, CRM197-PCV7 was administered together with a trivalent inactivated influenza vaccine (TIV) ([Jansen 2008](#)).

Four different interventions were used in five trials: OMPC-PCV7 in [Kilpi 2003](#) (a subset of these children received PPV23 as a booster dose); CRM197-PCV9 in [Dagan 2001](#); PHiD-CV10 in [Tregnaghi 2014](#)/[Sáez-Llorens 2017](#) and [Vesikari 2016](#)/[Karppinen 2019](#); and PHiD-CV11 in [Prymula 2006](#).

## Comparator

Control vaccines were used as comparators in all trials. The comparator vaccine in three trials was meningococcus type C conjugate vaccine (10 µg of group C oligosaccharide conjugated to carrier protein CRM197; MenC) ([Black 2000](#)/[Fireman 2003](#); [Dagan 2001](#); [O'Brien 2008](#)), whilst hepatitis A or B vaccine was used in the remaining eight trials.

## Outcome measures

Adverse effects were reported in nine trials including a total of 77,389 children ([Black 2000](#)/[Fireman 2003](#); [Dagan 2001](#); [Eskola 2001](#)/[Palmu 2009](#); [Jansen 2008](#); [Kilpi 2003](#); [Prymula 2006](#); [Tregnaghi 2014](#)/[Sáez-Llorens 2017](#); [Veenhoven 2003](#); [Vesikari 2016](#)/[Karppinen 2019](#)). [Tregnaghi 2014](#)/[Sáez-Llorens 2017](#) was part of the Clinical Otitis Media and Pneumonia Study (COMPAS; [clinicaltrials.gov/show/NCT00466947](https://clinicaltrials.gov/show/NCT00466947)), which assessed the efficacy and safety of PHiD-CV10 against invasive pneumococcal disease, community-acquired pneumonia, and AOM in 23,821 young Latin American children. Acute otitis media was studied in the Panama cohort only, which included 7357 children, whereas safety data were available for all 23,821 children.

Six trials applied a standardised diagnosis of AOM ([Eskola 2001](#)/[Palmu 2009](#); [Kilpi 2003](#); [Prymula 2006](#); [Tregnaghi 2014](#)/[Sáez-Llorens 2017](#); [van Kempen 2006](#); [Veenhoven 2003](#)), and one trial used standardised AOM registration forms to be completed by general practitioners ([Jansen 2008](#)). In two trials, AOM episodes were extracted from a computerised data source containing all visits registered by physicians ([Black 2000](#)/[Fireman 2003](#); [O'Brien 2008](#)). Two trials relied on parent-reported AOM episodes ([Dagan 2001](#); [Vesikari 2016](#)); [Vesikari 2016](#) used parent-reported, physician-confirmed AOM as the outcome of interest. [Karppinen 2019](#) used parent-reported data (symptom diaries), data from study clinic visits, and data on hospitalisations from an electronic registry as outcome measures. Two trials assessed outcomes during influenza seasons ([Jansen 2008](#); [van Kempen 2006](#)).

Seven trials also assessed the effect of PCVs on (serotype-specific) pneumococcal AOM ([Black 2000](#)/[Fireman 2003](#); [Eskola 2001](#); [Kilpi 2003](#); [O'Brien 2008](#); [Prymula 2006](#); [Tregnaghi 2014](#)/[Sáez-Llorens 2017](#); [Veenhoven 2003](#)). Three studies cultured middle ear fluid from all AOM episodes ([Eskola 2001](#); [Kilpi 2003](#); [Prymula 2006](#)), and one trial cultured middle ear fluid by tympanocentesis when fluid was suspected in the middle ear ([Tregnaghi 2014](#)/[Sáez-Llorens 2017](#)). One trial only cultured middle ear fluid from the first AOM episode by tympanocentesis or from spontaneously draining ears ([Veenhoven 2003](#)). Two trials assessed the effect on reported cultures that were obtained from spontaneously draining ears ([Black 2000](#)/[Fireman 2003](#); [O'Brien 2008](#)).

Three trials reported the effects of PCVs on recurrent AOM ([Black 2000](#)/[Fireman 2003](#); [Eskola 2001](#); [Prymula 2006](#)). Three studies included all types of otitis media, including, but not exclusively AOM, as an outcome ([Black 2000](#)/[Fireman 2003](#); [Dagan 2001](#); [O'Brien 2008](#)).

## Funding and conflicts of interest

Six trials were funded by pharmaceutical companies ([Black 2000](#)/[Fireman](#); [Eskola 2001](#)/[Palmu 2009](#); [Kilpi 2003](#); [Prymula 2006](#); [Tregnaghi 2014](#)/[Sáez-Llorens 2017](#); [Vesikari 2016](#)/[Karppinen 2019](#)). Three trials reported receiving support from non-

commercial (governmental) sources, but study vaccines were supplied by pharmaceutical companies (Jansen 2008; van Kempen 2006; Veenhoven 2003). One trial was supported by both a pharmaceutical company and governmental funding (O'Brien 2008). One trial reported that study vaccines were supplied by a pharmaceutical company (Dagan 2001).

#### Brief overview of clinical heterogeneity across included studies

There was considerable clinical heterogeneity across the included trials. There were differences in the timing of PCV administration, that is trials administering PCV during infancy and trials administering PCV later in life. As such, study populations varied from healthy infants to those at high risk of AOM. Secondly, the number of pneumococcal serotypes present in the vaccines, the type of conjugate method used, and co-administration of other vaccines differed substantially across trials. Study designs also varied, including both individually randomised controlled trials and cluster-RCTs. Finally, large differences in outcome assessments and AOM definitions were observed, varying from 'passive' (chart review at the end of the trial) to 'active' (parents were instructed to visit a physician in case of AOM symptoms) outcome assessments and physician-confirmed AOM episodes versus parent-reported AOM episodes. Consequently, AOM incidence in the control groups varied widely across the studies administering PCV during infancy,

from 0.13 to 1.3 episodes per person-year. We therefore did not perform meta-analyses.

#### Excluded studies

In the 2014 version of this review (Fortanier 2014), four studies were excluded because they (i) did not include a control vaccine (Gisselsson-Solen 2011); (ii) did not report outcome data relevant for this review (Jokinen 2012); (iii) assessed the effect of PCV on otitis media with effusion rather than AOM (Le 2007); and (iv) reported the effect of PCV on suppurative otitis media in an abstract of a conference meeting (Roy 2011). In the 2019 update, a further five studies were excluded that did not report outcome data relevant for this review (Palmu 2014; Palmu 2015a; Palmu 2015b; Palmu 2018; Sarasoja 2013). We did not exclude any new studies in this 2020 update. See [Characteristics of excluded studies](#) table.

#### Ongoing studies

We did not identify any ongoing studies relevant to this review.

#### Risk of bias in included studies

We judged the methodological quality of the included studies to be moderate to high. The 'Risk of bias' assessment is presented graphically in [Figure 2](#) and [Figure 3](#). See [Characteristics of included studies](#) table.

**Figure 2. 'Risk of bias' graph: review authors' judgements about each risk of bias item presented as percentages across all included studies.**

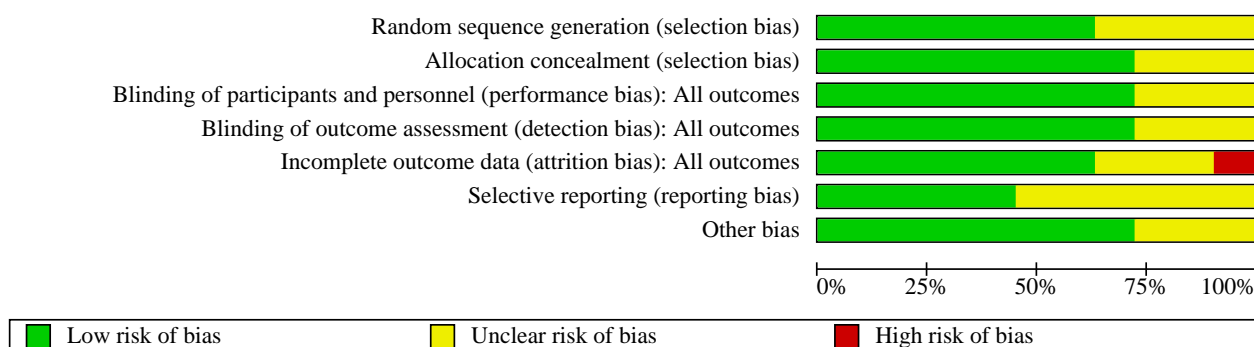


Figure 3. 'Risk of bias' summary: review authors' judgements about each risk of bias item for each included study.

	Random sequence generation (selection bias)	Allocation concealment (selection bias)	Blinding of participants and personnel (performance bias): All outcomes	Blinding of outcome assessment (detection bias): All outcomes	Incomplete outcome data (attrition bias): All outcomes	Selective reporting (reporting bias)	Other bias
Black 2000	?	?	?	+	?	?	?
Dagan 2001	?	+	+	?	+	?	+
Eskola 2001	+	+	+	+	+	?	+
Jansen 2008	?	?	+	+	?	+	+
Kilpi 2003	+	+	+	+	+	?	?
O'Brien 2008	+	+	+	?	?	+	?
Prymula 2006	+	+	?	+	+	+	+
Tregnaghi 2014	+	+	+	+	+	+	+
van Kempen 2006	?	+	+	+	+	?	+
Veenhoven 2003	+	?	+	+	+	?	+
Vesikari 2016	+	+	?	?	-	+	+



## Allocation

Eight trials described concealment of allocation adequately, whilst due to insufficient information we assessed this domain as unclear for three trials ([Black 2000](#)/[Fireman 2003](#); [Jansen 2008](#); [Veenhoven 2003](#)). We judged random sequence generation to be adequate in seven trials, whilst four trials provided insufficient information on methods of random sequence generation used ([Black 2000](#)/[Fireman 2003](#); [Dagan 2001](#); [Jansen 2008](#); [van Kempen 2006](#)).

## Blinding

Although all studies indicated that trials were double-blinded, three trials provided insufficient information about how blinding was performed ([Black 2000](#)/[Fireman 2003](#); [Prymula 2006](#); [Vesikari 2016](#)/[Karppinen 2019](#)).

## Incomplete outcome data

We judged risk of attrition bias to be high in one trial ([Vesikari 2016](#)/[Karppinen 2019](#)), unclear in three trials ([Black 2000](#)/[Fireman 2003](#); [Jansen 2008](#); [O'Brien 2008](#)), and low in seven trials.

## Selective reporting

We judged risk of reporting bias to be unclear in six trials, [Black 2000](#)/[Fireman 2003](#); [Dagan 2001](#); [Eskola 2001](#)/[Palmu 2009](#); [Kilpi 2003](#); [van Kempen 2006](#); [Veenhoven 2003](#), and low in the remaining five trials.

## Other potential sources of bias

We judged risk of bias due to other sources (including balances in baseline characteristics, use of co-intervention across groups, presence of formal sample size calculations, and (prespecified) interim analyses) as unclear in three trials, [Black 2000](#)/[Fireman 2003](#); [Kilpi 2003](#); [O'Brien 2008](#), and low in the remaining eight trials.

## Effects of interventions

See: [Summary of findings 1](#) *Pneumococcal conjugate vaccine versus control vaccine for preventing acute otitis media in children*

Effect estimates of the various PCV types, stratified by the age at which PCVs were administered and the occurrence of previous AOM/respiratory tract infection (RTI) episodes (i.e. administration in early infancy versus later in life), on frequency of all-cause AOM, (vaccine-type) frequency of pneumococcal AOM, and frequency of recurrent AOM (defined as three or more AOM episodes in six months or four or more in one year), are summarised in [Table 1](#), [Table 2](#), and [Table 3](#), respectively. The main results for PCVs administered in early infancy are described in [Summary of findings 1](#).

We included a total of 15 publications of 11 RCTs (60,733 children, range 74 to 37,868 per trial) of 7- to 11-valent PCVs versus control vaccines. Seven trials included infants who predominantly received primary vaccinations before six months of age (59,415 children in total) ([Black 2000](#)/[Fireman 2003](#); [Eskola 2001](#)/[Palmu 2009](#); [Kilpi 2003](#); [O'Brien 2008](#); [Prymula 2006](#); [Tregnaghi 2014](#)/[Sáez-Llorens 2017](#); [Vesikari 2016](#)/[Karppinen 2019](#)). One study included daycare attendees aged 12 to 35 months (264 children) ([Dagan 2001](#)). Two trials included children aged one to seven years with a history of AOM (457 children) ([van Kempen 2006](#); [Veenhoven 2003](#)). One trial included children aged 18 to 72 months with a previously diagnosed RTI (597 children) ([Jansen 2008](#)).

We have presented the results of individual trials as reported in the published papers; meta-analysis was inappropriate due to substantial differences amongst studies. We have assessed the statistical methods used to analyse data in each study.

## Adverse effects (co-primary outcome)

An overview of adverse effects reported in the individual studies can be found in [Table 4](#).

Mild local reactions and fever were common in both groups, occurring more frequently in the PCV than in the control vaccine groups: redness (< 2.5 cm): 5% to 20% versus 0% to 16%; swelling (< 2.5 cm): 5% to 12% versus 0% to 8%; and fever (< 39 °C): 15% to 44% versus 8% to 25%. More severe redness (> 2.5 cm), swelling (> 2.5 cm), and fever (> 39 °C) occurred less frequently (0% to 0.9%, 0.1% to 1.3%, and 0.4% to 2.5%, respectively) in children receiving PCV and did not differ significantly between PCV and control vaccine groups. Pain or tenderness, or both, was reported more frequently in children receiving PCV than in those receiving control vaccines: 3% to 38% versus 0% to 8%. Serious adverse events (SAEs) judged to be causally related to vaccination were rare and did not differ significantly between vaccine groups. No fatal SAE judged to be causally related to vaccination was reported.

The certainty of evidence for this outcome was high.

## Acute otitis media outcomes (co-primary outcome and secondary outcomes)

Seven studies used the generalised Cox proportional hazard method proposed by [Andersen 1982](#), currently regarded as the most optimal for analysing this kind of data ([Black 2000](#)/[Fireman 2003](#); [Eskola 2001](#)/[Palmu 2009](#); [Kilpi 2003](#); [Prymula 2006](#); [Tregnaghi 2014](#)/[Sáez-Llorens 2017](#); [van Kempen 2006](#); [Veenhoven 2003](#)).

[Dagan 2001](#) compared rates of AOM, but rather than comparing them by Poisson or negative binomial regression analysis (which would presumably yield results similar to those obtained with the Andersen approach), the Chi<sup>2</sup> test was used, which is suboptimal for comparing rates.

[Jansen 2008](#) used Poisson, and [Vesikari 2016](#)/[Karppinen 2019](#) used negative binomial regression analysis to compare rates of AOM between groups, accounting for the potential dependency of observations between individuals.

[O'Brien 2008](#) was a cluster-RCT that calculated incidence rate ratios with a Poisson regression with sandwich variance estimation to account for within-community correlation.

## Effect of PCV administered in early infancy (predominantly < 6 months of age)

Seven trials (59,415 children) included infants who predominantly received various types of PCV before six months of age ([Black 2000](#)/[Fireman 2003](#); [Eskola 2001](#)/[Palmu 2009](#); [Kilpi 2003](#); [O'Brien 2008](#); [Prymula 2006](#); [Tregnaghi 2014](#)/[Sáez-Llorens 2017](#); [Vesikari 2016](#)/[Karppinen 2019](#)).

## PCV7

In two trials (39,530 children), CRM197-PCV7 was the intervention for healthy infants aged two months ([Black 2000](#)/[Fireman 2003](#); [Eskola 2001](#)/[Palmu 2009](#)). The same vaccine was used as the intervention in one trial including 944 Navajo and White Mountain

Apache children aged up to two years. These children carry one of the highest risks of developing AOM in the world (O'Brien 2008).

In one trial (1666 children), OMPC-PCV7, with a subset of children receiving PPV23 as a booster dose, was used as the intervention in healthy infants aged two months (Kilpi 2003).

### Primary outcome

- Frequency of all-cause AOM episodes

In one trial including 37,868 healthy infants aged two months, CRM197-PCV7 was associated with a 6% (95% confidence interval (CI) 4% to 9%) relative risk reduction (RRR) in all-cause AOM episodes in an intention-to-treat (ITT) analysis (Black 2000/Fireman 2003). Per-protocol analysis of a trial including 1662 healthy infants aged two months showed that this same vaccine was associated with a non-significant 6% (95% CI -4% to 16%) RRR in all-cause AOM episodes (Eskola 2001/Palmu 2009).

In young children who carry a high baseline risk of developing AOM, CRM197-PCV7 was not associated with a reduction in all-cause AOM episodes (1 trial; 944 children; RRR -5%, 95% CI -25% to 12%; ITT analysis) (O'Brien 2008).

In one trial including 1666 healthy infants aged two months, OMPC-PCV7 was not associated with a reduction in all-cause AOM episodes in per-protocol analysis (RRR -1%, 95% CI -12% to 10%) (Kilpi 2003).

The certainty of evidence for the use of CRM197-PCV7 in low-risk infants for this outcome was moderate, downgraded one level due to imprecise effect estimate and study limitations (risk of bias). The certainty of evidence for the use of CRM197-PCV7 in young children with a high baseline risk of developing AOM and the use of OMPC-PCV7 in low-risk infants for this outcome was low; in both cases, the certainty of evidence was downgraded two levels due to the very imprecise effect estimate.

### Secondary outcomes

- Frequency of pneumococcal AOM

In one trial including 1662 healthy infants aged two months, CRM197-PCV7 was associated with a 20% (95% CI 7% to 31%) to 34% (95% CI 21% to 45%) RRR in pneumococcal AOM episodes in per-protocol analysis, depending on whether this outcome was assessed by a composite of positive culture or positive pneumolysin polymerase chain reaction (PCR) or by positive culture only (Eskola 2001/Palmu 2009).

In one trial including 1666 healthy infants aged two months, OMPC-PCV7 was associated with a 25% (95% CI 11% to 37%) RRR in pneumococcal AOM episodes in per-protocol analysis (Kilpi 2003).

The certainty of evidence for this outcome was high.

- Frequency of pneumococcal serotype-specific AOM

In two trials (39,530 healthy infants aged two months), administration of CRM197-PCV7 was associated with a 54% (95% CI 41% to 64%) to 65% (P = 0.04) RRR in vaccine-type pneumococcal AOM episodes in ITT analysis (Black 2000/Fireman 2003; Eskola 2001/Palmu 2009).

In one of these trials, CRM197-PCV7 was associated with a 51% (95% CI 27% to 67%) RRR in AOM episodes caused by cross-reactive serotypes and a non-significant 33% (95% CI -80% to 1%) relative increase in the risk of non-vaccine-type AOM episodes in per-protocol analyses (Eskola 2001/Palmu 2009).

In one trial (944 children), administration of CRM197-PCV7 in young children who carried a high baseline risk of developing AOM was associated with a non-significant 64% (95% CI -34% to 90%) RRR in vaccine-type pneumococcal AOM episodes in ITT analysis (O'Brien 2008).

In one trial including 1666 healthy infants aged two months, OMPC-PCV7 was associated with a 56% (95% CI 44% to 66%) RRR in vaccine-type pneumococcal AOM episodes in per-protocol analysis (Kilpi 2003). In the same trial, OMPC-PCV7 failed to show cross-protection (RRR -5%, 95% CI -47% to 25%), and this vaccine was associated with a non-significant 27% (95% CI -70% to 6%) relative increase in the risk of non-vaccine-type AOM episodes in per-protocol analyses (Kilpi 2003).

The certainty of evidence for the use of CRM197-PCV7 and OMPC-PCV7 in healthy infants for this outcome was high. However, the certainty of evidence for the use of CRM197-PCV7 in young children with high baseline risk of developing AOM for this outcome was moderate, downgraded one level due to study limitations (risk of bias) and imprecise effect estimate.

- Frequency of recurrent AOM

In two trials (39,530 children), administration of CRM197-PCV7 in healthy infants aged two months was associated with a 9% (95% CI -12% to 27%) to 10% (95% CI 7% to 13%) RRR in developing recurrent AOM (Black 2000/Fireman 2003; Eskola 2001/Palmu 2009).

The certainty of evidence for this outcome was moderate, downgraded one level due to imprecise effect estimate.

### PHiD-CV10/11

PHiD-CV10 was used as the intervention in two trials (12,307 children) (Tregnaghi 2014/Sáez-Llorens 2017; Vesikari 2016/Karppinen 2019). PHiD-CV11 was used in one trial (4968 children) (Prymula 2006).

### Primary outcome

- Frequency of all-cause AOM episodes

In one trial including 7359 healthy infants aged 6 to 16 weeks, PHiD-CV10 was associated with a non-significant 15% (95% CI -1% to 28%) RRR in all-cause AOM episodes in ITT analysis (Tregnaghi 2014/Sáez-Llorens 2017). Per-protocol analysis of a trial including 5095 healthy infants aged 6 weeks to 18 months showed that this same vaccine was associated with a non-significant 6% (95% CI -6% to 17%) RRR in all-cause AOM episodes (Vesikari 2016). In Karppinen 2019, a subcohort of 424 children nested within the FinIP trial, PHiD-CV10 was associated with a significant 23% (95% CI 0% to 40%) RRR in all-cause episodes of respiratory tract infections with AOM in per-protocol analysis.

In one trial including 4968 healthy infants aged 6 weeks to 5 months, PHiD-CV11 was associated with a 34% (95% CI 21% to 44%)

RRR in all-cause AOM episodes in per-protocol analysis (Prymula 2006).

However, it should be noted that the AOM incidence rates in the two trials with the largest point estimates, Prymula 2006; Tregnaghi 2014/Sáez-Llorens 2017, were low (Table 1). Consequently, the absolute risk differences in these trials were rather small.

The certainty of evidence for the use of PHiD-CV10 for this outcome was low, downgraded two levels due to study limitations (risk of bias) and imprecise effect estimates. The certainty of evidence for the use of PHiD-CV11 for this outcome was moderate, downgraded one level due to indirectness of evidence (low AOM incidence rate in the control group compared to other studies, most likely due to methodological differences with other studies).

#### Secondary outcomes

- Frequency of pneumococcal AOM

In one trial including 7359 healthy infants aged 6 to 16 weeks, PHiD-CV10 was associated with a 53% (95% CI 16% to 74%) RRR in pneumococcal AOM episodes in ITT analysis (Tregnaghi 2014/Sáez-Llorens 2017).

In one trial including 4968 healthy infants aged 6 weeks to 5 months, PHiD-CV11 was associated with a 52% (95% CI 37% to 63%) RRR in pneumococcal AOM episodes in per-protocol analysis (Prymula 2006).

The certainty of evidence for pneumococcal AOM episodes was high.

- Frequency of pneumococcal serotype-specific AOM

In one trial including 7359 healthy infants aged 6 to 16 weeks, PHiD-CV10 was associated with a 70% (95% CI 30% to 87%) RRR in vaccine-type pneumococcal AOM episodes in ITT analysis (Tregnaghi 2014/Sáez-Llorens 2017). In the same trial, PHiD-CV10 was associated with a non-significant 29% (95% CI -123% to 77%) RRR in AOM episodes caused by cross-reactive serotypes and a non-significant 15% (95% CI -153% to 71%) RRR in non-vaccine-type AOM episodes in ITT analyses (Tregnaghi 2014/Sáez-Llorens 2017).

In one trial including 4968 healthy infants aged 6 weeks to 5 months, PHiD-CV11 was associated with a 58% (95% CI 41% to 69%) RRR in vaccine-type pneumococcal AOM episodes in per-protocol analysis (Prymula 2006). In the same trial, PHiD-CV11 was associated with a 66% (95% CI 22% to 85%) RRR in AOM episodes caused by cross-reactive serotypes and a non-significant 9% (95% CI -64% to 49%) RRR in non-vaccine-type AOM episodes in per-protocol analyses (Prymula 2006).

The certainty of evidence for vaccine-type pneumococcal AOM episodes was high. The certainty of evidence for cross-reactive serotypes and non-vaccine-type AOM episodes was low, downgraded two levels due to very imprecise effect estimates.

- Frequency of recurrent AOM

In one trial including 4968 healthy infants aged 6 weeks to 5 months, PHiD-CV11 was associated with a non-significant 56% (95% CI -2% to 80%) RRR in developing recurrent AOM in per-protocol analysis (Prymula 2006).

The certainty of evidence for this outcome was low, downgraded two levels due to imprecise effect estimate and indirectness of evidence (low AOM incidence rate in control group compared to other studies, most likely due to methodological differences with other studies).

#### Effect of PCV administered at a later age (one year and above)

In three trials, various types of PCV7 were administered in children with a history of either RTI (597 participants), Jansen 2008, or AOM (457 participants in total) (van Kempen 2006; Veenhoven 2003).

#### CRM197-PCV7

##### Primary outcome

- Frequency of all-cause AOM episodes

In two trials (457 children) (van Kempen 2006; Veenhoven 2003), CRM197-PCV7 followed by PPV23 in children aged one to seven years with a history of AOM was not associated with further reductions in AOM episodes (1 trial; 383 children; RRR -25%, 95% CI -57% to 1%; ITT analysis (Veenhoven 2003); 1 trial; 74 children; RRR -16%, 95% CI -96% to 31%; per-protocol analysis (van Kempen 2006)).

In one trial including 597 children with a history of RTI, CRM197-PCV7 administered together with a trivalent influenza vaccine (CRM197-PCV7/TIV) was associated a 57% (95% CI 6% to 80%) RRR in all-cause AOM episodes compared to hepatitis B/placebo vaccination in per-protocol analysis (Jansen 2008). However, the effect of TIV/placebo compared to hepatitis B/placebo vaccination on all-cause AOM episodes appeared to be even larger (RRR 71%, 95% CI 30% to 88%) (Jansen 2008).

The certainty of evidence for this outcome was moderate, downgraded one level due to imprecise effect estimates.

##### Secondary outcomes

- Frequency of pneumococcal AOM

In per-protocol analysis of one trial including 383 children with a history of AOM, CRM197-PCV7 followed by PPV23 was associated with a non-significant 34% (P = 0.22) RRR in pneumococcal AOM episodes (Veenhoven 2003).

The certainty of evidence for this outcome was low, downgraded two levels due to very imprecise effect estimates (one study with a relatively small sample size).

- Frequency of pneumococcal serotype-specific AOM

In a per-protocol analysis of one trial including 383 children with a history of AOM, CRM197-PCV7 followed by PPV23 was associated with a non-significant 52% (P = 0.21) and 21% (P = 0.21) RRR in pneumococcal serotype-specific AOM and non-vaccine-type AOM episodes (Veenhoven 2003).

The certainty of evidence for this outcome was low, downgraded two levels due to very imprecise effect estimates (one study with a relatively small sample size).

- Frequency of recurrent AOM

None of the three trials in older children reported the effect of PCV7 on recurrent AOM.

## CRM197-PCV9

In one trial (264 children), CRM197-PCV9 was administered in healthy daycare attendees aged 12 to 35 months (Dagan 2001).

### Primary outcome

- Frequency of all-cause AOM episodes

In a per-protocol analysis, CRM197-PCV9 was associated with a non-significant 17% (95% CI -2% to 33%) RRR in all-cause otitis (OM) episodes (Dagan 2001).

The certainty of evidence for this outcome was very low, downgraded three levels due to study limitations (risk of bias and questions about outcome assessment) and imprecise effect estimate (one study with a relatively small sample size).

### Secondary outcomes

Dagan 2001 did not report on any of our secondary outcomes of interest.

## DISCUSSION

### Summary of main results

The current evidence base for the effects of PCVs in preventing AOM in children comes from 11 RCTs (60,733 children) of 7- to 11-valent PCVs versus control vaccines (meningococcus type C conjugate vaccine in three trials and hepatitis A or B vaccine in eight trials) with a generally low risk of bias. No relevant RCTs with the newer 13-valent PCV were available. In seven trials (59,415 children), PCVs were predominantly administered in child's first months of life, whilst four trials (1318 children) included children aged one year and over who were either healthy or who had a history of respiratory illness. There was considerable clinical heterogeneity across studies in terms of design, study population, type of PCV used, and outcome measures, therefore we did not perform meta-analyses.

The licenced CRM197-PCV7 and PHiD-CV10 vaccines, when administered during early infancy (< 6 months of age), were associated with substantial RRR in pneumococcal AOM (high-certainty evidence). However, their effects on all-cause AOM are far more uncertain, as most trials failed to demonstrate statistically significant differences for this outcome. Relative risk reductions for CRM197-PCV7 varied from -5% (95% CI -25% to 12%) in high-risk infants (low-certainty evidence) to 6% (95% CI -4% to 16%) and 6% (95% CI 4% to 9%) in low-risk infants (moderate-certainty evidence), whereas RRRs for PHiD-CV10 varied from 6% (95% CI -6% to 17%) to 15% (95% CI -1% to 28%) in healthy infants (low-certainty evidence).

Administration of PCVs in high-risk infants, after early infancy, and in older children with a history of respiratory illness or frequent AOM was not associated with reductions in all-cause AOM.

Local redness, swelling, fever, and tenderness/pain were commonly reported and occurred more frequently in children receiving PCV than in those receiving control vaccines, but these adverse effects were mostly mild. More severe redness (> 2.5 cm), swelling (> 2.5 cm), and fever (> 39 °C) occurred far less frequently and did not differ between vaccine groups. Serious adverse events judged to be causally related to vaccination were rare and did not

differ significantly between vaccine groups. No fatal SAE judged to be causally related to vaccination was reported.

### Overall completeness and applicability of evidence

The 11 RCTs included in this review differed substantially in terms of RCT type, study population (age of PCV administration and AOM baseline risk), PCV type (vaccine valency, carrier protein, and booster regimen), co-administration of other vaccines, and AOM assessment and definition used. Furthermore, in the infant studies focusing on AOM bacteriology (Eskola 2001; Kilpi 2003; Prymula 2006; Tregnaighi 2014), the control groups varied markedly in the proportions of *Streptococcus pneumoniae*, *Haemophilus influenzae*, and *Moraxella catarrhalis* found in the middle ear fluid. This might be related to time and geographic region as well as case definition, and has important implications for the effects of PCV on preventing all-cause AOM episodes. Additionally, three studies included older otitis-prone children, so the intervention was aimed at secondary or even tertiary prevention, and not primary prevention (Jansen 2008; van Kempen 2006; Veenhoven 2003). The reduced efficacy of CRM197-PCV7 in children with a history of AOM may be explained by an increased susceptibility to subsequent infections, not only with non-vaccine-type pneumococci, but also other nasopharyngeal colonisers, due to 'damage' already suffered by the middle ear mucosa caused by prior AOM (Veenhoven 2003). Another explanation, whilst debated, could be the non-protective, impaired antibody responses of children who are otitis-prone (Pichichero 2013; Wiertsema 2012). It thus appears that the age at which PCV is administered, a history of AOM episodes, or both, modifies the effect of PCV on AOM, despite the fact that age alone could not be identified as a statistically significant effect modifier (Black 2000/Fireman 2003; Veenhoven 2003). Our review did not focus on the effects of PCVs on shifts in serotypes over time. Further research into the impact of PCVs on (serotype) replacement is warranted, since a shift in causative pathogens may have considerable implications for both AOM burden and vaccine effectiveness.

### Quality of the evidence

The certainty of evidence varied substantially per outcome measure. For the frequency of all-cause AOM and recurrent AOM, the certainty of evidence varied from moderate (further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate) to low (further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate). The certainty of evidence for adverse effects and frequency of pneumococcal AOM was high (further research is very unlikely to change our confidence in the estimate of effect).

### Potential biases in the review process

We adhered to the prespecified review protocol. Two review authors (JLHdS, RPV for the 2020 update) independently searched all relevant electronic databases using a search syntax comprising all relevant synonyms for PCV and AOM. We also performed a broad internet search to identify potentially relevant articles. To increase the yield of relevant studies, we reviewed the reference lists of all identified studies and systematic reviews or meta-analyses. We searched ClinicalTrials.gov and the WHO ICTRP for completed and ongoing trials. For the 2020 update, a new review author (JLHdS) independently reviewed 'Risk of bias' and GRADE assessments and



data extraction of all included studies. Any discrepancies with the findings described in the 2019 update were discussed with a second review author (RPV), and where needed, with a third review author (RAMJD).

## Agreements and disagreements with other studies or reviews

Our main findings are in agreement with three other systematic reviews on the effect of PCV in children, indicating that PCVs provide substantial protection against pneumococcal AOM, but that their effects on all-cause AOM are more uncertain and far less pronounced (Ewald 2016; Pavia 2009; Taylor 2012).

In AOM, there is a high potential for replacement by other bacterial pathogens that are common colonisers of the nasopharynx. CRM197-PCV7 is known to affect nasopharyngeal carriage of pneumococci, with a shift from vaccine-type pneumococci to non-vaccine-type pneumococci and other otopathogens including non-typeable *H influenza* and *Staphylococcus aureus* (Biesbroek 2014; Block 2006; Casey 2013; Coker 2010; Eskola 2001; Obaro 1996; Somech 2011; van Gils 2011; Wiertsema 2011). Nasopharyngeal carriage results from a recent RCT on PHiD-CV10 showed similar bacterial colonisation patterns as observed in CRM197-PCV7 amongst healthy Dutch children aged up to two years (van den Bergh 2013). The middle ear is directly connected to the nasopharynx, and by lowering the carriage of vaccine-type pneumococci, a niche may be created for other bacteria with pathogenic potential (Block 2006; Veenhoven 2003; Veenhoven 2004). Recent studies have shown that nationwide implementation of PCVs may have changed the frequency of the causative otopathogens involved in AOM towards pneumococcal serotypes not included in the vaccines and non-typeable *H influenza* (Ben-Shimol 2019; Casey 2013; Coker 2010; Kaur 2017; Somech 2011; Wiertsema 2011).

Although RCT data failed to demonstrate a convincing beneficial effect of CRM197-PCV7 and PHiD-CV10 on all-cause AOM, various global postmarketing studies with these licenced vaccines, as well as the newer CRM197-PCV13, suggest that the impact of PCVs on AOM may be substantial (Eythorsson 2018; Gisselsson-Solen 2017; Kawai 2018; Lau 2015; Lecrenier 2020; Magnus 2012; Marom 2014; Poehling 2007; Sigurðsson 2018; Soysal 2020; Zhou 2008), which may be attributable to indirect (herd) effects of vaccination. However, it should be noted that findings from observational studies warrant careful interpretation, as variability in baseline incidence, study population, and case definition, as well as fluctuations in risk factors for AOM such as breastfeeding, household smoking, daycare attendance rates, and implementation of AOM clinical practice guidelines, may affect the AOM incidences reported. For example, results from Boston (USA) showed that the decline in uncomplicated AOM, treatment failure, and AOM relapse was at least as large in the 2000-to-2004 period compared to the 1996-to-2000 period, leaving the 'true' contribution of PCV in reducing AOM incidence uncertain (Sox 2008). Furthermore, reduced exposure to household smoking amongst other factors such as PCV7 coverage since 2002, may have contributed to the steady decline in USA paediatric ambulatory visits for otitis media over the period of 1993 to 2006 (Alpert 2011).

The impact of PHiD-CVs may expand beyond their effects on pneumococcal AOM to AOM caused by non-typeable *H influenza* due to the carrier protein D (Forsgren 2008). A recent

review including pre-clinical, clinical, and postmarketing studies concluded that PHiD-CVs may decrease AOM caused by non-typeable *H influenza*, but that more evidence including pathogen-specific outcomes is clearly warranted (Clarke 2017). The diversity of non-typeable *H influenza*, with some strains lacking protein D, may limit the effect of PHiD-CVs on non-typeable *H influenza* AOM. In our review, administration of the licenced PHiD-CV10 in healthy infants was associated with non-significant 6% (95% CI -6% to 17%) and 15% (95% CI -1% to 28%) relative reductions in the risk of all-cause AOM. The added benefit of PHiD-CV10 over the previously licenced CRM197-PCV7 on all-cause AOM therefore remains uncertain.

We found limited evidence that administration of PCVs during infancy may reduce the risk of recurrent AOM. This is in line with accumulating evidence that PCVs might disrupt the continuum of evolution from pneumococcal-associated otitis media towards chronic/recurrent otitis media by prevention of early vaccine-serotype AOM, thereby reducing subsequent and more complex disease caused by non-vaccine serotypes and non-typeable *H influenza* (Ben-Shimol 2014; Dagan 2016). These findings are further supported by secondary analyses of some of the trials included in our review indicating that licenced CRM197-PCV7 and PHiD-PCV10 lead to fewer ventilation tubes insertions for chronic/recurrent otitis media (Black 2000; Palmu 2015b; Sarasoja 2013).

## AUTHORS' CONCLUSIONS

### Implications for practice

The current evidence from randomised controlled trials (RCTs) indicates that, albeit associated with large relative risk reductions in pneumococcal acute otitis media (AOM), the effect of administration of the licenced CRM197-PCV7 and PHiD-CV10 in healthy, low-risk infants on AOM is derived from low- to moderate-certainty evidence and is therefore far more uncertain. In addition, we found no evidence of a beneficial effect on all-cause AOM of administering PCVs in high-risk infants, after early infancy, and in older children with a history of respiratory illness.

However, global postmarketing studies of these vaccines and of the licenced CRM197-PCV13 suggest that the impact (i.e. both direct and indirect effects) of pneumococcal conjugate vaccines (PCVs) administered in infancy on AOM may be substantial. Furthermore, it should be noted that the decision of whether or not to implement PCV should not come from AOM studies only, but also from studies that see and show the big picture, including data on invasive pneumococcal disease such as pneumonia, bacteraemia, and meningitis.

Compared to control vaccines, PCVs were associated with an increase in mild local reactions (redness, swelling), fever, and pain/tenderness. However, we found no evidence of a difference in (far less frequently occurring) more severe local reactions, fever, or serious adverse events judged to be causally related to vaccination.

### Implications for research

Since most countries across the world have implemented PCV in nationwide immunisation programmes, future RCTs comparing PCVs versus control vaccines are unlikely to be performed. Whilst there is some observational evidence of a difference in effects on AOM between PHiD-CV10 and the newer CRM197-PCV13 (Gisselsson-Solen 2017), future trials may compare the efficacy

of various types of PCVs. More importantly, future research will likely shed a light on the effects of other vaccines to prevent AOM, including (protein-based) vaccines directed at *Streptococcus pneumoniae* (Hammitt 2019), as well as other pathogens including non-typeable *Haemophilus influenzae* and *Moraxella catarrhalis* (Pettigrew 2017).

Whether any decline in AOM will continue or wane over time due to replacement is relevant and deserves ongoing monitoring. Besides a reduction of nasopharyngeal vaccine-type serotypes, which is presumed to induce herd effects, replacing pneumococcal serotypes may not only lead to replacement disease in vaccines, but also in the population. Continuing surveillance of nasopharyngeal carriage and pneumococcal disease in both the short and long term

(Spijkerman 2012), in different settings and geographic locations, is therefore of the utmost importance.

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\* Indicates the major publication for the study

## CHARACTERISTICS OF STUDIES

### Characteristics of included studies [ordered by study ID]

#### Black 2000

##### Study characteristics

Methods	<p><b>Randomised:</b> yes, at individual level</p> <p><b>Design:</b> standard parallel-group design</p> <p><b>Intention-to-treat:</b> yes</p> <p><b>Follow-up:</b> 6 to 31 months</p>
Participants	<p><b>N:</b> 37,868 healthy infants</p> <p><b>Age:</b> 2 months</p> <p><b>Setting:</b> 23 medical centres within Northern California Kaiser Permanente (NCKP), USA</p> <p><b>Inclusion criteria:</b> healthy children aged 2 months</p> <p><b>Exclusion criteria:</b> children with sickle cell disease, known immunodeficiency, any serious chronic or progressive disease, a history of seizures, or a history of either pneumococcal or meningococcal disease</p> <p><b>Baseline characteristics:</b> not described</p>
Interventions	<p>Children were randomly allocated to either CRM197-PCV7 or a meningococcus type C conjugate vaccine (10 µg of group C oligosaccharide conjugated to carrier protein CRM197; MenC) at 2, 4, 6 and 12 to 15 months of age</p> <p><b>Tx:</b> CRM197-PCV7; N = 18,927 received 1 dose or more of the vaccine (unclear how many children were included in otitis media analyses)</p>

## Black 2000 (Continued)

**C:** MenC; N = 18,941 received 1 dose or more of the vaccine (unclear how many children were included in otitis media analyses)

**Additional vaccines:** routine childhood vaccines were administered at the recommended ages: DTwP or DTaP; oral poliovirus vaccine or inactivated poliovirus vaccine; Hib; hepatitis B; measles-mumps-rubella vaccine; varicella.

Initially all participants received a vaccine combining *Haemophilus b* conjugate and DTwP into the opposite leg and oral poliovirus vaccine concurrently. When recommendations changed, the protocol was amended to allow administration of DTaP and inactivated poliovirus vaccine. Vaccines not given concomitantly were given at least 2 weeks apart from study vaccine.

Outcomes	<p><b>Primary outcome:</b> invasive pneumococcal disease caused by vaccine serotypes</p> <p><b>Secondary outcomes:</b> number of otitis media episodes in fully vaccinated per protocol; number of otitis media visits; time to recurrent otitis media (defined as 3 or more episodes in 6 months or 4 or more in 12 months); number of tympanostomy tubes placements; number of cases of spontaneously draining ruptured tympanic membranes with culture of a vaccine serotype pneumococcus; safety (local and systemic reactions at 48 to 72 hours after vaccination, uncommon events requiring medical attention 30 days and 60 days after vaccination, and mortality)</p> <p>Clinical diagnoses of AOM were obtained from computerised data sources using diagnoses registered by emergency physicians and paediatricians in the NCKP population. Each clinic visit constituted a new episode unless it was classified as a follow-up visit. A visit &lt; 21 days after another otitis media visit was always considered a follow-up visit. A visit 42 days or more after the most recent otitis media visit was considered a new episode. Visits occurring between 21 and 42 days, if the appointment was made &lt; 3 days in advance, were considered new episodes.</p>	
Funding sources	The study was supported by an unrestricted grant from Wyeth(-Ayerst); authors were employed at Kaiser Permanente Vaccine Study Center (Oakland), Wyeth Lederle Vaccines and Pediatrics (Pearl River), University of Pennsylvania (Philadelphia), and Vanderbilt University Medical Center (Nashville).	
Declarations of interest	Not described; authors were employed at Kaiser Permanente Vaccine Study Center (Oakland), Wyeth Lederle Vaccines and Pediatrics (Pearl River), University of Pennsylvania (Philadelphia), and Vanderbilt University Medical Center (Nashville)	
Notes		
<b>Risk of bias</b>		
<b>Bias</b>	<b>Authors' judgement</b>	<b>Support for judgement</b>
Random sequence generation (selection bias)	Unclear risk	Method of random sequence generation not described
Allocation concealment (selection bias)	Unclear risk	No method of allocation concealment described
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Indicated as a double-blind study, but insufficient details provided to ensure blinding of participants and personnel
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Clinical diagnoses of AOM were obtained from computerised data sources using diagnoses registered by emergency physicians and paediatricians (non-trialists).
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Unclear how many children were included in otitis media analyses

## Black 2000 (Continued)

Selective reporting (reporting bias)	Unclear risk	Study protocol is not available. Otitis media endpoint (efficacy against otitis media episodes) is reported as a secondary endpoint.
Other bias	Unclear risk	Study enrolment was stopped as a result of prespecified interim analysis.

## Dagan 2001

### Study characteristics

Methods	<p><b>Randomised:</b> yes, at individual level</p> <p><b>Design:</b> standard parallel-group design</p> <p><b>Intention-to-treat:</b> no, per-protocol analysis</p> <p><b>Follow-up:</b> 2 years starting 1 month after complete immunisation</p>
Participants	<p><b>N:</b> 264 healthy infants (261 children were included in clinical follow-up)</p> <p><b>Age:</b> 12 to 35 months</p> <p><b>Setting:</b> 8 daycare centres in Beer-Sheva, Israel</p> <p><b>Inclusion criteria:</b> healthy children aged 12 to 35 months</p> <p><b>Exclusion criteria:</b> children who had received any vaccine within the previous 4-week period, or who were scheduled to receive any vaccine during the 4 weeks after the administration of the study vaccines, or who had received immunoglobulin within 8 weeks of study vaccination; known or suspected impairment of immunologic functions; major congenital malformation or serious chronic disease; known hypersensitivity to any components of the study vaccine; previous severe vaccine-associated adverse reaction; previous vaccination with any pneumococcal or meningococcal vaccine; febrile illness (rectal temperature 38 °C) within 72 h before vaccination</p> <p><b>Baseline characteristics:</b> described and balanced (Table 1 of trial publication)</p>
Interventions	<p>Children were randomly allocated to either CRM197-PCV9 or MenC. Children aged 12 to 17 months at time of enrolment received 2 intramuscular injections 2 to 3 months apart, whilst those 18 to 35 months at time of enrolment received 1 intramuscular injection.</p> <p><b>Tx:</b> CRM-197-PCV9; N = 131</p> <p><b>C:</b> MenC; N = 130</p> <p><b>Additional vaccines:</b> not described</p>
Outcomes	<p><b>Primary outcome:</b> nasopharyngeal carriage of <i>Streptococcus pneumoniae</i> of the serotypes found in the vaccines in general and antibiotic-resistant <i>S pneumoniae</i> in particular</p> <p><b>Secondary outcomes:</b> parent-reported respiratory infections including otitis media, tolerance (tenderness, local and systemic reactions after vaccination including erythema, induration, and fever)</p> <p>18 encounters were planned for each child during the 2-year follow-up period. Encounters were planned to take place monthly during the first year and bimonthly during the second year. At each visit the parents were questioned about illness and antibiotic use since the last visit. Illness episodes were divided into 4 categories:</p> <p>(1) upper respiratory infections; (2) lower respiratory problems; (3) otitis media; and (4) other illnesses. Only episodes starting 1 month after complete immunisation were counted.</p>
Funding sources	The study vaccines were provided by Wyeth-Lederle Vaccines and Pediatrics (Pearl River, NY).
Declarations of interest	Not described



## Dagan 2001 (Continued)

Notes

**Participants lost to follow-up during first 12 months:total:** 32/261 (12.3%)

**Participants lost to follow-up during first 12 months:Tx:** 16/131 (12.2%)

**Participants lost to follow-up during first 12 months:C:** 16/130 (12.3%)

### Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Method of random sequence generation not described. Block randomisation (n = 6) stratified by DCC and age.
Allocation concealment (selection bias)	Low risk	Randomisation list provided in a sealed envelope by Wyeth-Lederle Vaccines.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	PCV9 and MenC vaccines dissimilar in appearance. 2 nurses not belonging to the study team injected the vaccines. They were not allowed to reveal the child's allocation.
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Parental interview. A positive report of OM was defined as an episode.
Incomplete outcome data (attrition bias) All outcomes	Low risk	Follow-up rates reported in Table 1. 12% of children followed up for < 12 months.
Selective reporting (reporting bias)	Unclear risk	Study protocol is not available.
Other bias	Low risk	No other sources of bias identified.

## Eskola 2001

### Study characteristics

Methods	<p>This trial was part of a study including <a href="#">Kilpi 2003</a> (FinOM Vaccine Trial). Both <a href="#">Eskola 2001</a> and <a href="#">Kilpi 2003</a> used the same control group (hepatitis B vaccine containing 5 µg of recombinant hepatitis B surface protein) but a different treatment group, each with a different PCV7 type. <a href="#">Eskola 2001</a> used CRM197-PCV7, whilst <a href="#">Kilpi 2003</a> used OMPC-PCV7.</p> <p><b>Randomised:</b> yes, at individual level</p> <p><b>Design:</b> standard parallel-group design</p> <p><b>Intention-to-treat:</b> yes, both ITT and per-protocol analysis described</p> <p><b>Follow-up:</b> 22 consecutive months (children were followed up to 24 months of age)</p>
Participants	<p><b>N:</b> 1662 healthy infants</p> <p><b>Age:</b> 2 months</p> <p><b>Setting:</b> 8 study clinics in the communities of Tampere, Kangasala, and Nokia, Finland</p> <p><b>Inclusion criteria:</b> healthy children aged 2 months</p>

**Eskola 2001** (Continued)

**Exclusion criteria:** not described

**Baseline characteristics:** described and balanced (Table 1 of trial publication)

Interventions	<p>Children were randomly allocated to either CRM197-PCV or a hepatitis B vaccine at 2, 4, 6 and 12 to 15 months of age.</p> <p><b>Tx:</b> CRM197-PCV; N = 831 (N = 786 completed the follow-up as specified in the protocol)  <b>C:</b> hepatitis B vaccine; N = 831 (N = 794 completed the follow-up as specified in the protocol)  <b>Additional vaccines:</b> a combination vaccine containing whole-cell DTP and Hib was given in the child's opposite thigh at the same visit as the pneumococcal vaccine at 2, 4, and 6 months of age. In half of the study clinics, the carrier protein in the DTP and <i>Haemophilus influenzae</i> vaccine was CRM197, whilst in the other half it was tetanus toxoid. Inactivated poliovirus vaccine was given at 7 months of age and again at the same time as the fourth dose of the study vaccine at 12 months of age. Measles–mumps–rubella vaccine was administered at 18 months.</p>
Outcomes	<p><b>Primary outcome:</b> number of AOM episodes due to the pneumococcal serotypes included in the vaccine</p> <p><b>Secondary outcomes:</b> number of all-cause AOM episodes, culture-confirmed and pathogen-specific AOM episodes; preventing first and subsequent AOM episodes; number of children with recurrent AOM episodes (defined as 3 or more AOM episodes in the last 6 months or 4 or more in the last 12 months); serious adverse events, safety (pain, local and systemic reactions within 3 days after vaccination, unexpected events after vaccination and mortality)</p> <p>All children attended 1 of the study clinics for enrolment at 2 months of age and thereafter at 4, 6, 7, 12, 13, 18, and 24 months. Parents were encouraged to bring their child to the study clinic for evaluation of symptoms suggesting respiratory infection or AOM. AOM was diagnosed by otoscopy (visibly abnormal tympanic membrane in terms of colour, position, or mobility, suggesting middle ear effusion) and the presence of at least 1 of the following symptoms or signs of acute infection: fever, earache, irritability, diarrhoea, vomiting, acute otorrhoea not caused by otitis externa, and other symptoms of respiratory infection.</p> <p>For the overall and pathogen-specific AOM episodes, a new episode was considered to have started if at least 30 days had elapsed since the beginning of the previous episode. For AOM episodes according to serotype, a new episode was considered to have started if 30 days had elapsed since the beginning of an episode due to the same serotype, or if any interval had elapsed since the beginning of an episode due to a different serotype. If more than 1 serotype was recovered from the middle ear fluid at the same time, only 1 episode was considered to have started.</p>
Funding sources	Supported by Merck, Pasteur Mérieux Connaught, and Wyeth-Lederle Vaccines and Pediatrics
Declarations of interest	Dr Eskola and Dr Kilpi have served as consultants to Wyeth-Lederle Vaccines.
Notes	<p><b>Participants lost to follow-up:total:</b> 82/1662 (4.9%) did not complete the follow-up period specified in the protocol</p> <p><b>Participants lost to follow-up:Tx:</b> 45/831 (5.4%) did not complete the follow-up period specified in the protocol</p> <p><b>Participants lost to follow-up:C:</b> 37/831 (4.5%) did not complete the follow-up period specified in the protocol</p>
<b>Risk of bias</b>	
<b>Bias</b>	<b>Authors' judgement</b> <b>Support for judgement</b>
Random sequence generation (selection bias)	<p>Low risk</p> <p>6 letters corresponding to the 3 treatment options were randomly allocated to consecutive participant identification numbers, using an allocation of 1:1:1 and a block size of 12 (see Kilpi 2003).</p>



**Eskola 2001** (Continued)

Allocation concealment (selection bias)	Low risk	Individual treatment assignments were kept in sealed envelopes until vaccination (see <a href="#">Kilpi 2003</a> ).
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Use of vaccinators who were not otherwise involved in the trial follow-up. Letter code was destroyed immediately after vaccination (see <a href="#">Kilpi 2003</a> ).
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Assessment of the outcome was done according to a strict definition of AOM. Assessment was performed by personnel other than those who vaccinated the children (vaccinators were not otherwise involved in the trial follow-up) (see <a href="#">Kilpi 2003</a> ).
Incomplete outcome data (attrition bias) All outcomes	Low risk	Reasons for dropout or lost to follow-up, or both, were not reported. This is not expected to have a major impact on outcome since 94.6% in the CRM197-PCV7 and 95.5% in the control group completed the follow-up as specified in the protocol.
Selective reporting (reporting bias)	Unclear risk	Prespecified outcomes (primary and secondary) are listed in ClinicalTrials.gov (although uploaded after study end).
Other bias	Low risk	No other sources of bias identified.

**Jansen 2008**
**Study characteristics**

Methods	<p><b>Randomised:</b> yes, at individual level</p> <p><b>Design:</b> standard parallel-group design</p> <p><b>Intention-to-treat:</b> yes</p> <p><b>Follow-up:</b> follow-up started 14 days after the second set of vaccinations and continued for 6 to 18 months, depending on the year of inclusion</p>
Participants	<p><b>N:</b> 597 children with a previously diagnosed RTI</p> <p><b>Age:</b> 18 to 72 months</p> <p><b>Setting:</b> GPs in the centre of the Netherlands selected children</p> <p><b>Inclusion criteria:</b> children aged 18 to 72 months with a previously diagnosed RTI registered according to the ICPC, i.e. AOM; cough (with fever); acute upper RTI; acute laryngitis/tracheitis; acute bronchitis/bronchiolitis; influenza; pneumonia; pleurisy/pleural effusion</p> <p><b>Exclusion criteria:</b> children with chronic asthma or recurrent wheezing (for longer than 3 months) treated with corticosteroids; craniofacial abnormalities; clinically significant hypersensitivity to eggs; previous serious adverse reactions to vaccines; previous influenza, pneumococcal, or hepatitis B vaccinations and those with conditions for which these vaccinations are already recommended, such as chronic cardiac and respiratory conditions</p> <p><b>Baseline characteristics:</b> described and balanced (Table 1 of trial publication)</p>
Interventions	<p>Children were randomly allocated to either TIV/PCV7, TIV/placebo (TIV plus standard diluent (0.9% phosphate buffered NaCl)), or HBV/placebo (recombinant HBV vaccine-Engerix B Junior plus placebo vaccine). Strains in the TIV 2003-2004 formulation were H1N1, H3N2, and B/HongKong/330/01; strains in the TIV 2004-2005 formulation were H1N1, H3N2, and B/Shanghai/361/2002; strains in the TIV 2005-2006 formulation included H1N1, H3N2, and B/Shanghai/361/2002.</p>

**Jansen 2008** (Continued)

Children received 2 vaccinations 4 to 8 weeks apart in the first year of inclusion; the first 2 cohorts of children received a subsequent vaccination in the subsequent year.

**Tx:** TIV/CRM197-PCV7; N = 197 (N = 163 completed; 67,867 person-days analysed, 14% missing)

**C1:** TIV/placebo; N = 187 (N = 148 completed; 60,515 person-days analysed, 20% missing)

**C2:** HBV/placebo; N = 195 (N = 160 completed; 67,679 person-days analysed, 15% missing)

**Additional vaccines:** not described

Outcomes	<p><b>Primary outcome:</b> febrile RTI, defined as fever (tympanic temperature 38.0 °C) for at least 2 consecutive days accompanied by 1 or more of the aforementioned signs or symptoms of RTI with a moderate or severe severity score</p> <p><b>Secondary outcomes:</b> febrile RTI-related PCR-confirmed influenza, GP visits, antibiotic prescriptions, or a physician-diagnosed episode of AOM, tolerability and safety.</p> <p>Each parent was instructed to keep a daily diary in which they recorded any clinical signs or symptoms associated with RTI and characterised their severity on a scale of 1 (mild) to 3 (severe). The parent was also instructed to measure the child's body temperature using a validated electronic tympanic thermometer, and was asked to record all GP visits due to their child's RTI-related complaints. For each such visit, the GP was instructed to complete a form including information on the diagnosis and possible antibiotic prescriptions.</p> <p>During influenza seasons, the parent was instructed to contact the trial centre for evaluation for influenza if the child had fever (tympanic temperature 38.0 °C) for more than 1 day accompanied by at least 1 RTI-associated sign or symptom of severity score 2. A trained research assistant obtained a nasopharyngeal swab for viral determination within 4 days of onset of fever and symptoms. Each sample was analysed by real-time PCR for the presence of influenza A and B viruses.</p>
Funding sources	<p>The study was funded by the Netherlands Organisation for Health Research and Development (Zon-Mw). The funding agency played no role in the design and conduct of the study; the collection, analysis, and interpretation of the data; or the preparation, review, or approval of the manuscript.</p> <p>Influenza vaccines were provided by Solvay, Weesp, the Netherlands. Pneumococcal vaccines were provided by Wyeth Vaccines Research, Berkshire, UK. Hepatitis B vaccines were provided by GlaxoSmithKline BV, Rixensart, Belgium.</p>
Declarations of interest	The authors declared no conflicts of interest.
Notes	<p><b>Participants lost to follow-up:total:</b> 108/579 (18.7%) completely (N = 41) or partially (N = 67) lost to follow-up</p> <p><b>Participants lost to follow-up:Tx:</b> 34/197 (17.3%) completely (N = 8) or partially (N = 26) lost to follow-up; 67,867 person-days analysed, 14% missing</p> <p><b>Participants lost to follow-up:C1:</b> 39/187 (20.8%) completely (N = 19) or partially (N = 20) lost to follow-up; 60,515 person-days analysed, 20% missing</p> <p><b>Participants lost to follow-up:C2:</b> 35/195 (17.9%) completely (N = 14) or partially (N = 21) lost to follow-up; 67,679 person-days analysed, 15% missing</p> <p>2 of the 3 treatment arms received an additional vaccination in the second year of the study. To evaluate blinding, parents of these cohorts of children were asked which vaccinations they thought their child had received just after the vaccinations were given and at the end of the study. Just after the vaccination, 87% of the parents either did not know or identified the wrong set of vaccinations; at the end of the study, this percentage was 80%, indicating successful blinding.</p>
<b>Risk of bias</b>	
<b>Bias</b>	<b>Authors' judgement      Support for judgement</b>

**Jansen 2008** (Continued)

Random sequence generation (selection bias)	Unclear risk	Method of random sequence generation not described; children were randomly assigned in blocks of 3 in a 1:1:1 ratio.
Allocation concealment (selection bias)	Unclear risk	No method of allocation concealment was described.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	The injections were administered by non-blinded research nurses who were not involved in subsequent follow-up and who were instructed not to reveal the intervention allocation. The treatment group assignments were not revealed to parents, investigators, research personnel conducting the follow-up, or healthcare providers, all of whom remained blinded throughout the study.
Blinding of outcome assessment (detection bias) All outcomes	Low risk	The parents were asked to record all GP visits due to their child's RTI-related complaints. For each such visit, the GP was instructed to complete a form including information on the diagnosis and possible antibiotic prescriptions. The treatment group assignments were not revealed to parents, investigators, and research personnel conducting the follow-up, or healthcare providers, all of whom remained blinded throughout the study.
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Substantial loss to follow-up (< 14% in both groups)
Selective reporting (reporting bias)	Low risk	Prespecified outcomes (primary and secondary) are listed in ClinicalTrials.gov.
Other bias	Low risk	No other sources of bias identified.

**Kilpi 2003**
**Study characteristics**

Methods	<p>This trial was part of a study including <a href="#">Eskola 2001</a> (FinOM Vaccine Trial). Both <a href="#">Eskola 2001</a> and <a href="#">Kilpi 2003</a> used the same control group (hepatitis B vaccine containing 5 µg of recombinant hepatitis B surface protein) but a different PCV7 type. <a href="#">Eskola 2001</a> used CRM197-PCV7, whilst <a href="#">Kilpi 2003</a> used OM-PC-PCV7.</p> <p><b>Randomised:</b> yes, at individual level</p> <p><b>Design:</b> standard parallel-group design</p> <p><b>Intention-to-treat:</b> no, per-protocol analysis</p> <p><b>Follow-up:</b> 22 consecutive months (children were followed up to 24 months of age)</p>
Participants	<p><b>N:</b> 1666 healthy infants</p> <p><b>Age:</b> 2 months</p> <p><b>Setting:</b> 8 study clinics in the communities of Tampere, Kangasala, and Nokia, Finland</p> <p><b>Inclusion criteria:</b> healthy children aged 2 months</p> <p><b>Exclusion criteria:</b> not described</p> <p><b>Baseline characteristics:</b> described and balanced (Table 1)</p>
Interventions	Children were randomly allocated to either OMPC-PCV7 or a hepatitis B vaccine at 2, 4, 6 and 12 to 15 months of age. From 3 November 1997 onwards, for the children randomised to receive OMPC-PCV7,

## Kilpi 2003 (Continued)

the fourth dose of the conjugate vaccine was replaced by PPV23, Pneumovax23 including serotypes 1, 2, 3, 4, 5, 6B, 7F, 8, 9N, 9V, 10A, 11A, 12F, 14, 15B, 17F, 18C, 19F, 19A, 20, 22F, 23F, and 33F.

**Tx:** OMPC-PCV7; N = 835 (N = 805 completed the follow-up as specified in the protocol)

**C:** hepatitis B vaccine; N = 831 (N = 794 completed the follow-up as specified in the protocol)

**Additional vaccines:** a diphtheria–tetanus toxoids–pertussis vaccine with a whole-cell pertussis component, combined with a *Haemophilus influenzae* type b conjugate vaccine (DTP-Hib), was administered concomitantly with the first 3 doses of the study vaccine; an inactivated poliovirus vaccine was administered with the fourth dose. In 4 study clinics, the carrier protein in the DTP-Hib conjugate combination was CRM197, whilst in the other 4 study clinics it was tetanus toxoid.

Outcomes	See <a href="#">Eskola 2001</a> .
Funding sources	Supported by Aventis Pasteur, Merck, and Wyeth-Lederle Vaccines and Pediatrics
Declarations of interest	Not described; from <a href="#">Eskola 2001</a> : Dr Eskola and Dr Kilpi have served as consultants to Wyeth-Lederle Vaccines
Notes	<p><b>Participants lost to follow-up:total:</b> 67/1666 (4.0%) did not complete the follow-up period specified in the protocol</p> <p><b>Participants lost to follow-up:Tx:</b> 30/835 (3.6%) did not complete the follow-up period specified in the protocol</p> <p><b>Participants lost to follow-up:C:</b> 37/831 (4.5%) did not complete the follow-up period specified in the protocol</p>

### Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	6 letters corresponding to the 3 treatment options were randomly allocated to consecutive participant identification numbers, using an allocation of 1:1:1 and a block size of 12.
Allocation concealment (selection bias)	Low risk	Individual treatment assignments were kept in sealed envelopes until vaccination.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Use of vaccinators who were not otherwise involved in the trial follow-up. Letter code was destroyed immediately after vaccination.
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Assessment of the outcome was done according to a strict definition of AOM. Assessment was performed by personnel other than those who vaccinated the children (vaccinators were not otherwise involved in the trial follow-up).
Incomplete outcome data (attrition bias) All outcomes	Low risk	No reporting of reasons for dropout or loss to follow-up. This is not expected to have a major impact on outcome since 96.0% in the OMPC-PCV7 and 95.5% in the control group completed the follow-up as specified in the protocol.
Selective reporting (reporting bias)	Unclear risk	Prespecified outcomes (primary and secondary) are listed in ClinicalTrials.gov (although uploaded after study end).
Other bias	Unclear risk	Mixed schedule with 187 children boosted with PPV23. Unclear how researchers identified those allocated to OMPC-PCV7 to receive PPV23 after November 1997

## O'Brien 2008

### Study characteristics

Methods	<p>The design of this cluster-randomised trial has been described extensively in <a href="#">Moulton 2001</a>, whilst the findings on IPD (main outcome of the trial) are published in <a href="#">O'Brien 2003</a>.</p> <p><b>Randomised:</b> yes, at group level</p> <p><b>Design:</b> cluster-randomised design</p> <p><b>Intention-to-treat:</b> no, per-protocol analysis</p> <p><b>Follow-up:</b> depending on time of inclusion, maximum duration of follow-up 40 months</p>
Participants	<p><b>N:</b> 944 (944 of the 4476 children were randomly selected for chart review. This sample size was determined by logistic feasibility and expected frequency of healthcare events. Of the 944 children, 856 were found to have strictly met the chart review criteria.)</p> <p><b>Age:</b> below 2 years</p> <p><b>Setting:</b> Navajo and White Mountain Apache region, USA</p> <p><b>Inclusion criteria:</b> Navajo and White Mountain Apache children below 2 years of age</p> <p><b>Exclusion criteria:</b> no exclusion criteria described</p> <p><b>Baseline characteristics:</b> balanced, but data not shown</p>
Interventions	<p>Children were randomly allocated to either CRM197-PCV7 or MenC (10 µg of group C oligosaccharide conjugated to carrier protein CRM197). For each of the study and control vaccines, 3 immunisation schedules were designed according to age of entry into the trial: 6 weeks to 6 months (3 doses, ideally at 2, 4, and 6 months of age and a booster at 12 to 15 months of age), 7 months to 11 months (2 doses 1 month apart and a booster at 12 to 15 months of age), and 12 months to 23 months (2 doses separated by at least 2 months). Over the course of the trial, the great majority of new enrollees were in the first group, which is referred to as the primary efficacy cohort.</p> <p><b>Tx:</b> CRM197-PCV7; N = unknown (N = 424 analysed in primary efficacy group)</p> <p><b>C:</b> MenC; N = unknown (N = 432 analysed in primary efficacy group)</p> <p><b>Additional vaccines:</b> not described</p>
Outcomes	<p><b>Primary outcome:</b> clinically diagnosed episodes of OM</p> <p>Every medical visit made by study children was evaluated through 2 years of age. OM visits, as documented by the children's treating physician, were recorded.</p> <p>A new OM episode was counted if any of the following were recorded as the diagnosis: OM, AOM, bilateral OM, chronic OM, OM with perforation, otorrhoea, pressure-equalising tube placement, perforated tympanic membrane, serous OM and bullous myringitis.</p> <p>An episode of AOM was categorised as either AOM or bilateral AOM. An OM episode was categorised as severe if there were 3 or more OM visits for the episode. A child's first medical visit for OM was considered their first episode. OM visits occurring fewer than 21 days after the immediately prior otitis-related visit and visits noted as a follow-up to a previous otitis-related visit were counted as follow-up visits, not as OM episodes.</p>
Funding sources	<p>Financial support for the American Indian PnCRM7 Efficacy Trial was from Wyeth Vaccines, the US National Institutes of Health, World Health Organization, the National Vaccine Program Office, and the Centers for Disease Control and Prevention.</p>
Declarations of interest	<p>Dr O'Brien and Dr Santosham participated in Wyeth Scientific Advisory Boards. Dr O'Brien, Dr Moulton, Dr Reid, Dr Weatherholtz, and Dr Santosham received research funding from Wyeth Vaccines.</p>
Notes	<p><b>Participants lost to follow-up:</b>total: 88/944 (9.3%) not included in primary efficacy analysis</p>

## O'Brien 2008 (Continued)

**Participants lost to follow-up:Tx:** unknown

**Participants lost to follow-up:C:** unknown

### Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Computer-generated randomisation using 38 independent randomisation units, stratified using 3 blocks of 4 units and 13 blocks of 2 units
Allocation concealment (selection bias)	Low risk	6 labels were assigned to the vaccines (B, F, H, M, T, U), with 3 labels for CRM197-PCV7 and 3 for MenC. The grouping of these codes was known only to a statistician employed by the manufacturer (who had no other responsibilities with respect to the trial other than handling treatment allocation and randomisation issues). No loss of clusters
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Masked treatment assignment (vaccines were labelled). In addition, field staff were blinded as to serotype of the invasive disease cases, and thus did not know which ones would be likely to be prevented by an effective vaccine.
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Every medical visit made by study children was evaluated through 2 years of age. OM visits, as documented by the children's treating physician, were recorded. Treating physicians were blinded to treatment allocation.
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	88 of the 944 children (9.3%) not included in primary efficacy analysis; no information provided on the distribution across treatment groups
Selective reporting (reporting bias)	Low risk	Study design was described extensively in <a href="#">Moulton 2001</a> and <a href="#">O'Brien 2003</a> .
Other bias	Unclear risk	Study enrolment was stopped as a result of prespecified interim analysis.

## Prymula 2006

### Study characteristics

Methods	<p><b>Randomised:</b> yes, at individual level</p> <p><b>Design:</b> standard parallel-group design</p> <p><b>Intention-to-treat:</b> yes, both ITT and per-protocol analysis described</p> <p><b>Follow-up:</b> efficacy follow-up started on the day of the first dose of study vaccine (for ITT analysis) or 2 weeks after the third vaccine dose (for the per-protocol analysis) and continued until 24 to 27 months of age</p>
Participants	<p><b>N:</b> 4968 healthy infants</p> <p><b>Age:</b> between 6 weeks and 5 months</p> <p><b>Setting:</b> 27 paediatric centres in the Czech Republic and 23 in Slovakia</p> <p><b>Inclusion criteria:</b> healthy children aged between 6 weeks and 5 months with no acute illness</p> <p><b>Exclusion criteria:</b> use of any investigational or non-registered drug or vaccine other than the study vaccines within 30 days preceding first dose of the study vaccines; previous vaccination against <i>Strep-</i></p>

## Prymula 2006 (Continued)

*tococcus pneumoniae*; fever (defined as a rectal temperature of 38 °C or higher or temperature by other routes of 37.5 °C or higher); history of allergic disease or reactions likely to be exacerbated by any component of the study vaccines; other conditions that might potentially interfere with the interpretation of study outcomes according to the investigator

**Baseline characteristics:** described and balanced (Table 1 of trial publication)

Interventions	<p>Children were randomly allocated to either PHiD-CV11 or a hepatitis A vaccine (containing 720 ELISA units of inactivated hepatitis A virus antigen (strain HM 175)) at about 3, 4, 5 and 12 to 15 months of age.</p> <p><b>Tx:</b> PHiD-CV11; N = 2489 (N = 2455 included in per-protocol cohort for efficacy)  <b>C:</b> hepatitis A vaccine; N = 2479 (N = 2452 included in per-protocol cohort for efficacy)  <b>Additional vaccines:</b> a concomitant hexavalent diphtheria-tetanus-3-component acellular pertussis-hepatitis B-inactivated poliovirus types 1, 2, and 3 <i>Haemophilus influenzae</i> type b (DTPa-HBV-IPV/Hib) vaccine was offered to all study participants, followed by a booster dose at age 15 to 18 months</p>
Outcomes	<p><b>Primary outcome:</b> first episode of AOM caused by vaccine pneumococcal serotypes</p> <p><b>Secondary outcomes:</b> first episode of AOM caused by non-typeable <i>H influenzae</i>, any all-cause AOM episodes, any vaccine-type AOM episodes, any cross-reactive serotypes AOM, any non-vaccine-type AOM, safety (adverse events arising within 31 days of vaccination, and serious adverse events occurring throughout the study period).</p> <p>There was no active surveillance. Unscheduled doctor visits could take place any time during follow-up according to standard local practice (parents consulting their local paediatrician in case of illness of their child). Parents were advised to consult their paediatrician if their child was sick, had ear pain, or had spontaneous ear discharge. Children with suspected AOM were immediately referred to ENT surgeons.</p> <p>AOM was defined as either abnormal findings of the tympanic membrane at otoscopy (i.e. redness, bulging, loss of light reflex) or the presence of middle ear effusion as shown by simple or pneumatic otoscopy or by microscopy together with at least 2 of the following signs or symptoms: ear pain, ear discharge, hearing loss, fever, lethargy, irritability, anorexia, vomiting, or diarrhoea. These signs or symptoms had to be present for a maximum of 14 days.</p> <p>For children with repeated doctor visits, a new episode of AOM was judged to have started if more than 30 days had elapsed since the beginning of the previous episode. Additionally, for categories defined according to bacterial pathogen or serotype, a new episode was judged to have started if any interval had elapsed since the beginning of an episode caused by a different bacterial pathogen or serotype.</p> <p>Recurrent AOM was defined as 3 or more AOM episodes in the last 6 months or 4 or more in the last 12 months.</p>
Funding sources	The study was supported by GlaxoSmithKline Biologicals, Rixensart, Belgium.
Declarations of interest	Dr Prymula is a consultant to GlaxoSmithKline and other pharmaceutical companies, and has received travel grants or honoraria paid by healthcare companies within the past 3 years. 7 co-authors are employees of GlaxoSmithKline Biologicals, of which 4 own shares in GlaxoSmithKline.
Notes	<p><b>Participants lost to follow-up:total:</b> 61/4968 (1.2%) did not complete the follow-up period specified in the protocol</p> <p><b>Participants lost to follow-up:Tx:</b> 34/2489 (1.4%) did not complete the follow-up period specified in the protocol</p> <p><b>Participants lost to follow-up:C:</b> 27/2479 (1.1%) did not complete the follow-up period specified in the protocol</p>
<b>Risk of bias</b>	
<b>Bias</b>	<b>Authors' judgement      Support for judgement</b>

## Prymula 2006 (Continued)

Random sequence generation (selection bias)	Low risk	Computer-generated random list
Allocation concealment (selection bias)	Low risk	Randomisation (1:1) was done with a study-specific central randomisation system via the internet which, on receipt of the infant's initials and birth date, determined the vaccine number to be used.
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Indicated as a double-blinded study. The sponsor numbered the vaccine supplies. However, it is unknown whether the appearance of the vaccines was similar at the time of administration.
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Visits during efficacy follow-up were according to standard local clinical practice. When AOM was suspected, children were referred to ENT surgeons.
Incomplete outcome data (attrition bias) All outcomes	Low risk	No reporting of reasons for dropout or loss to follow-up. This is not expected to have a major impact on outcome since 98.6% in the PHiD-CV11 and 98.9% in the control group completed the follow-up as specified in the protocol.
Selective reporting (reporting bias)	Low risk	Prespecified outcomes (primary and secondary) are listed in ClinicalTrials.gov.
Other bias	Low risk	Study enrolment was stopped as a result of prespecified interim analysis. No other sources of bias identified.

## Tregnaghi 2014

### Study characteristics

Methods	<p>This trial was part of Clinical Otitis Media and Pneumonia Study (COMPAS; <a href="https://clinicaltrials.gov/show/NCT00466947">clinicaltrials.gov/show/NCT00466947</a>) to assess the efficacy of PHiD-CV10 against IPD, CAP, and AOM in young Latin American children.</p> <p><b>Randomised:</b> yes, at individual level</p> <p><b>Design:</b> standard parallel-group design</p> <p><b>Intention-to-treat:</b> yes, both ITT and per-protocol analysis described</p> <p><b>Follow-up:</b> total follow-up duration 4 years</p>
Participants	<p><b>N:</b> 23,821 healthy infants (for Panama, AOM cohort: 7359)</p> <p><b>Age:</b> mean age 9 months</p> <p><b>Setting:</b> well-baby clinics at 5 sites (3 in Argentina, 1 in Colombia, and 1 in Panama); all countries are classified as upper-middle economies</p> <p><b>Inclusion criteria:</b> healthy children aged 6 to 16 weeks</p> <p><b>Exclusion criteria:</b> use or planned use of any investigational or unregistered drug or vaccine other than the study vaccines; previous vaccination against diphtheria, tetanus, pertussis, <i>Haemophilus influenzae</i> type b, hepatitis A, and/or <i>Streptococcus pneumoniae</i>; history of allergic disease or reactions likely to be exacerbated by any components of the study vaccines; acute disease at time of enrolment; low birthweight (&lt; 2500 g) not permitted for Colombia</p> <p><b>Baseline characteristics:</b> described and balanced (Table 3 of trial publication)</p>



**Tregnaghi 2014** (Continued)

Interventions	<p>Children were randomly allocated to either PHiD-CV10 or a hepatitis B vaccine at 2, 4, and 6 months followed by 1 dose of PHiD-CV10 or hepatitis A vaccine at 15 to 18 months of age.</p> <p><b>Tx:</b> PHiD-CV10; N = 11,875 (N = 10,295 completed the follow-up as specified in the protocol); for AOM cohort N = 3602 (N = 3010 completed the follow-up as specified in the protocol)</p> <p><b>C:</b> hepatitis B vaccine; N = 11,863 (N = 10,201 completed the follow-up as specified in the protocol); for AOM cohort N = 3612 (N = 2979 completed the follow-up as specified in the protocol)</p> <p><b>Additional vaccines:</b> a combination vaccine containing diphtheria-tetanus-acellular pertussis-inactivated polio and <i>H influenzae</i> type b (DTPa-IPV/Hib) was given in the child's opposite thigh at the same visit as the pneumococcal vaccine at 2, 4, 6 and 15 to 18 months</p>
Outcomes	<p><b>Primary outcome:</b> likely bacterial CAP</p> <p><b>Secondary outcomes:</b> other CAP outcomes, first episode of clinically confirmed AOM, first episode of pathogen-specific AOM, serious adverse events and mortality occurring throughout the study period.</p> <p>The AOM outcome was studied in Panama only (7357 of the 23,821 randomised children).</p> <p>Initially, AOM cases were captured only when parents sought medical attention for children with AOM symptoms. However, because of a lower-than-expected AOM rate, the surveillance was enhanced in July 2009 (2 years after start of enrolment) through regular telephone calls or home visits by study personnel who advised parents to visit the clinic if their child had symptoms suggestive of AOM. If the physician suspected AOM, the child was referred to 1 of the ENT surgeons involved in the trial.</p> <p>Clinically confirmed AOM was defined as either altered visual appearance of the tympanic membrane (e.g. redness, bulging, loss of light reflex) or the presence of middle ear effusion (by pneumatic otoscopy or otomicroscopy). Recent onset (duration of less than 5 days) of at least 2 of the following clinical symptoms was also required: ear pain, ear discharge, hearing loss, fever, lethargy, irritability, anorexia, vomiting, or diarrhoea.</p> <p>The severity of each AOM episode (mild, moderate, severe) was assessed by combining objective elements of the Friedman scale (the Ear Treatment Group-five items (ETG-5) and otoscopy scale with eight grades of severity (OS-8)) with subjective elements in a clinical otologic scale.</p> <p>When middle ear fluid was suspected, tympanocentesis was performed, and bacterial presence was assessed by culture.</p>
Funding sources	<p>Sponsored by GlaxoSmithKline Biologicals, the vaccine developer and manufacturer. The data generated in the trial are subject to a confidentiality agreement between the investigators and the sponsor that allowed the investigators full access to the study data and included an obligation for GlaxoSmithKline Biologicals to permit publication without excessive delay.</p>
Declarations of interest	<p>Dr Sáez-Llorens declares having received financial support from the study sponsor for travel to meetings, and his institution has received grants from Health Research International. Drs López and Calvo declare that their institutions have received support for travel to meetings and grants from the study sponsor. Dr Calvo declares that her institution has received consulting fee/honorary from the study sponsor. Dr Hausdorff is a patent coholder of PCV13 (no royalties). 9 co-authors are employees of GlaxoSmithKline companies and own stock/stock options from the GlaxoSmithKline group of companies.</p>
Notes	<p><b>Participants lost to follow-up:total:</b> 592/3602 (16.4%) did not complete the follow-up period specified in the protocol</p> <p><b>Participants lost to follow-up:Tx:</b> 633/3612 (17.5%) did not complete the follow-up period specified in the protocol</p> <p><b>Participants lost to follow-up:C:</b> 1225/7214 (17.0%) did not complete the follow-up period specified in the protocol</p>
<b>Risk of bias</b>	
<b>Bias</b>	<b>Authors' judgement      Support for judgement</b>

**Tregnaghi 2014** (Continued)

Random sequence generation (selection bias)	Low risk	Quote: "The randomization list was generated by the sponsor using a standard SAS (SAS Institute) program and was used to number the vaccines. A randomization blocking scheme was used to ensure that balance between treatment groups was maintained"
Allocation concealment (selection bias)	Low risk	Quote: "Vaccine allocation at each site was performed using a central randomization system on the Internet (SBIR, GlaxoSmithKline Vaccines), and treatment was concealed from all study personnel"
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Vaccines were numbered by the sponsor, and treatment allocation was concealed from study personnel. There were minor differences in vaccine appearance, but vaccines were prepared and administered by personnel who took no further part in the study.
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Vaccines were prepared and administered by personnel who took no further part in the study. Parents/guardians of participating children and study personnel involved in data gathering, processing, and analysis and safety assessment were blind to vaccine allocation.
Incomplete outcome data (attrition bias) All outcomes	Low risk	Almost all randomised children were included in ITT analysis.
Selective reporting (reporting bias)	Low risk	Prespecified outcomes listed at <a href="https://clinicaltrials.gov/show/NCT00466947">clinicaltrials.gov/show/NCT00466947</a> .
Other bias	Low risk	No other sources of bias identified.

**van Kempen 2006**
**Study characteristics**

Methods	<p>This study was performed in parallel with <a href="#">Veenhoven 2003</a> (OMAVAX-trial), but analysed separately due to differences in study population.</p> <p><b>Randomised:</b> yes, at individual level</p> <p><b>Design:</b> standard parallel-group design</p> <p><b>Intention-to-treat:</b> unclear</p> <p><b>Follow-up:</b> 26 months</p>
Participants	<p><b>N:</b> 74 children with a history of AOM</p> <p><b>Age:</b> between 1 and 7 years</p> <p><b>Setting:</b> ENT department of the Ghent University Hospital in Belgium</p> <p><b>Inclusion criteria:</b> children aged 1 to 7 years with a history of AOM defined as at least 2 separate clinically diagnosed AOM episodes in the past year</p> <p><b>Exclusion criteria:</b> children with any underlying illnesses including immunocompromising conditions other than partial serum IgA and IgG2 deficiencies, craniofacial abnormalities, previous pneumococcal vaccination, or documented hypersensitivity to any of the vaccine components</p> <p><b>Baseline characteristics:</b> described and balanced (Table 1 of trial publication)</p>

**van Kempen 2006** (Continued)

Interventions	<p>Children were randomly allocated to either a CRM197-PCV7 or a hepatitis A vaccine (containing 720 units of inactivated hepatitis A virus). Children aged 12 to 24 months received 2 intramuscular injections with a 1-month interval, whilst those aged over 2 years received 1 intramuscular injection. Children allocated to CRM197-PCV7 additionally received PPV23 6 months (in children aged 12 to 24 months) or 7 months (in those aged over 2 years) later.</p> <p><b>Tx:</b> CRM197-PCV7 plus PPV23; N = 38 (N = 35 completed the vaccination scheme) <b>C:</b> hepatitis A vaccine; N = 36 (N = 33 completed the vaccination scheme) <b>Additional vaccines:</b> not described</p>	
Outcomes	<p><b>Primary outcome:</b> number of AOM episodes during the 18-month follow-up</p> <p><b>Secondary outcomes:</b> immunogenicity; nasopharyngeal carriage of conjugate vaccine-related serotypes; and antibiotic-resistant pneumococci.</p> <p>At scheduled hospital visits at 7, 14, 20, and 26 months after randomisation, a medical history was taken, antibiotic usage noted, and an otomicroscopic examination performed. When at least 1 month following complete vaccination a new AOM episode was suspected, parents were asked to bring their sick child to the study centre within 24 hours for otoscopic diagnosis. In case of all other AOM episodes during follow-up, participants were allowed to visit the study centre, their family physician, or a paediatrician, who was asked to report otoscopic findings, diagnosis, and treatment on an AOM registration form.</p> <p>AOM was defined as an abnormal tympanic membrane on otomicroscopy (red, dull, or bulging) plus at least 1 of the following symptoms or signs of acute infection: earache, acute otorrhoea, fever (&gt; 38.5 °C rectally), or irritability.</p>	
Funding sources	<p>Study was supported by the Netherlands Organisation for Health Research and Development (ZonMw) and the Dutch health insurance company Zilveren Kruis-Achmea as part of the OMAVAX-trial. Wyeth-Lederle Vaccines and Pediatrics provided the pneumococcal vaccines, and GlaxoSmithKline provided the hepatitis A vaccines.</p>	
Declarations of interest	<p>The authors declared no conflicts of interest.</p>	
Notes	<p><b>Participants lost to follow-up:total:</b> 6/74 (8.1%) did not complete the follow-up period specified in the protocol</p> <p><b>Participants lost to follow-up:Tx:</b> 3/38 (7.9%) did not complete the follow-up period specified in the protocol</p> <p><b>Participants lost to follow-up:C:</b> 3/36 (8.3%) did not complete the follow-up period specified in the protocol</p>	
<b>Risk of bias</b>		
<b>Bias</b>	<b>Authors' judgement</b>	<b>Support for judgement</b>
Random sequence generation (selection bias)	Unclear risk	Method of random sequence generation not described, randomisation stratified according to age (12 to 24 months versus 25 to 84 months) and number of previous AOM episodes per year (2 to 3 versus 4 or more episodes).
Allocation concealment (selection bias)	Low risk	2 study nurses immunised all children according to a randomisation list provided to them in a sealed envelope by a third party (the Julius Center for Health Sciences, Utrecht, the Netherlands).
Blinding of participants and personnel (performance bias) All outcomes	Low risk	The nurses that vaccinated children were not allowed to reveal the child's allocation to either the study team or the parents.

## van Kempen 2006 (Continued)

Blinding of outcome assessment (detection bias) All outcomes	Low risk	When a new AOM episode was suspected, parents were asked to bring their sick child to the study centre within 24 hours for otoscopic diagnosis. In case of all other AOM episodes during follow-up, participants were allowed to visit the study centre, their family physician, or a paediatrician, who was asked to report otoscopic findings, diagnosis, and treatment on an AOM registration form.
Incomplete outcome data (attrition bias) All outcomes	Low risk	In total, 6 of the 74 children (8.1%) did not complete the follow-up period specified in the protocol (equally distributed across groups). Reasons for withdrawal are described in the Results section of the article.
Selective reporting (reporting bias)	Unclear risk	No study protocol available.
Other bias	Low risk	No other sources of bias identified.

## Veenhoven 2003

### Study characteristics

Methods	<p>This study was performed in parallel with <a href="#">van Kempen 2006</a>, but analysed separately due to differences in study population.</p> <p><b>Randomised:</b> yes, at individual level</p> <p><b>Design:</b> standard parallel-group design</p> <p><b>Intention-to-treat:</b> yes</p> <p><b>Follow-up:</b> 18 months, starting 1 month after completion of the vaccination scheme</p>
Participants	<p><b>N:</b> 383 children with a history of AOM</p> <p><b>Age:</b> between 1 and 7 years</p> <p><b>Setting:</b> a general hospital (Spaarne Hospital, Haarlem) and a tertiary care hospital (Wilhelmina Children's Hospital of the University Medical Centre Utrecht) in the Netherlands</p> <p><b>Inclusion criteria:</b> children aged 1 to 7 years with a history of AOM defined as 2 or more AOM episodes in the year before study entry. The number of previous AOM episodes was based on parental report and on clinical confirmation of the diagnosis by a physician.</p> <p><b>Exclusion criteria:</b> children with immunodeficiency, cystic fibrosis, immotile cilia syndrome, craniofacial abnormalities, chromosomal abnormalities such as Down's syndrome, and severe adverse events during previous vaccinations</p> <p><b>Baseline characteristics:</b> described and balanced (Table 1 of trial publication)</p>
Interventions	<p>Children were randomly allocated to either CRM197-PCV7 followed by a PPV23 or a hepatitis A or B vaccine.</p> <p>Children aged 12 to 24 months in the pneumococcal vaccination group received PCV7 twice with a 1-month interval, followed 6 months later by PPV23. The control vaccine group received 3 hepatitis B vaccinations (Engerix-B) according to a similar time schedule.</p> <p>Children aged 25 to 84 months in the pneumococcal vaccine group received 1 dose of PCV7 followed 7 months later by PPV23. The control group received hepatitis A vaccine (Havrix) twice.</p> <p><b>Tx:</b> CRM197-PCV7 plus PPV23; N = 190 (N = 190 included in ITT analysis)</p> <p><b>C:</b> hepatitis A or B vaccine; N = 193 (N = 193 included in ITT analysis)</p> <p><b>Additional vaccines:</b> not described</p>

**Veenhoven 2003** (Continued)

Outcomes	<p><b>Primary outcome:</b> number of clinical AOM episodes during the 18-month follow-up</p> <p><b>Secondary outcomes:</b> number of AOM episodes due to the 7 pneumococcal serotypes included in the PCV7 vaccine and nasopharyngeal carriage of conjugate vaccine serotypes, serious adverse events</p> <p>Parents were instructed to visit the study clinics or their GP, ENT surgeon, or paediatrician to assess symptoms suggesting AOM. Physicians registered signs and symptoms of every AOM episode on standard registration forms and were unaware of treatment allocation. AOM was defined according to the guideline issued by the Dutch College of General Practitioners, i.e. presence of an abnormal tympanic membrane on otoscopy (red, dull, or bulging) or otorrhoea and at least 1 of the following signs or symptoms of acute infection: acute earache, new-onset otorrhoea, irritability, or fever greater than 38.5 °C rectally or 38.0 °C axillary.</p>	
Funding sources	Study was supported by the Netherlands Organisation for Health Research and Development (ZonMw) and the Dutch health insurance company Zilveren Kruis-Achmea as part of the OMAVAX-trial. Wyeth-Lederle Vaccines and Pediatrics provided the pneumococcal vaccines, and GlaxoSmithKline provided the hepatitis A vaccines.	
Declarations of interest	The authors declared no conflicts of interest.	
Notes	<p><b>Participants lost to follow-up:total:</b> 1/383 (0.3%); all children included in ITT analysis</p> <p><b>Participants lost to follow-up:Tx:</b> 0/190 (0%)</p> <p><b>Participants lost to follow-up:C:</b> 1/193 (0.5%)</p>	
<b>Risk of bias</b>		
<b>Bias</b>	<b>Authors' judgement</b>	<b>Support for judgement</b>
Random sequence generation (selection bias)	Low risk	Table of random numbers that identified the vaccine scheme, randomisation stratified according to age (12 to 24 months versus 25 to 84 months) and number of previous AOM episodes per year (2 to 3 versus 4 or more episodes)
Allocation concealment (selection bias)	Unclear risk	No method of allocation concealment was described.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Vaccine was administered to the child by a study nurse, so that parents and physicians were unaware of treatment allocation.
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Parents were instructed to visit the study clinics or their family physician, otolaryngologist, or paediatrician to assess symptoms suggesting AOM. Physicians registered signs and symptoms of every AOM episode on standard registration forms and were unaware of treatment allocation. AOM was defined according to the guideline issued by the Dutch College of General Practitioners.
Incomplete outcome data (attrition bias) All outcomes	Low risk	All randomised children were included in ITT analysis.
Selective reporting (reporting bias)	Unclear risk	No study protocol available.
Other bias	Low risk	No other sources of bias identified.

## Vesikari 2016

### Study characteristics

Methods	<p>This trial was nested within the Finnish invasive pneumococcal disease vaccine trial (FinIP; <a href="https://clinicaltrials.gov/show/NCT00861380">clinicaltrials.gov/show/NCT00861380</a>; <a href="#">Palmu 2013</a>), a cluster-randomised double-blind trial to assess the efficacy of PHiD-CV10 against IPD, all-cause antibiotic purchases, tympanostomy tube placements, and vaccine-preventable diseases.</p> <p><b>Randomised:</b> yes, at cluster level</p> <p><b>Design:</b> parallel-group (4 groups) design</p> <p><b>Intention-to-treat:</b> no, per-protocol analysis</p> <p><b>Follow-up:</b> mean duration of follow-up 18 months</p>
Participants	<p><b>N:</b> 5095 healthy infants</p> <p><b>Age:</b> mean age at first dose 2.3 months</p> <p><b>Setting:</b> 15 study centres in Finland</p> <p><b>Inclusion criteria:</b> healthy children aged 6 weeks to 18 months</p> <p><b>Exclusion criteria:</b> prior administration of pneumococcal vaccine, hepatitis A or B vaccine, any investigational or non-registered product, contraindication to immunisation</p> <p><b>Baseline characteristics:</b> described and balanced (Suppl Table 1)</p>
Interventions	<p>Children were randomly allocated (2:2:1:1) to 1 of PHiD-CV10 3 + 1, 2 + 1, control 3 + 1, control 2 + 1 (control vaccine was hepatitis B for children aged &lt; 12 months or hepatitis A for those aged 12 months and above).</p> <p><b>Tx1:</b> PHiD-CV10 3 + 1; N = 1846 (N = 1846 completed the follow-up as specified in the protocol)</p> <p><b>Tx2:</b> PHiD-CV10 2 + 1; N = 1313 (N = 942 completed the follow-up as specified in the protocol)</p> <p><b>C1:</b> hepatitis A or B vaccine 3 + 1; N = 1073 (N = 468 completed the follow-up as specified in the protocol)</p> <p><b>C2:</b> hepatitis A or B vaccine 2 + 1; N = 861 (N = 861 completed the follow-up as specified in the protocol)</p> <p><b>Additional vaccines:</b> a combination vaccine containing diphtheria-tetanus-acellular pertussis-inactivated polio and <i>Haemophilus influenzae</i> type b (DTPa-IPV/Hib) and human rotavirus vaccine were given at the same visit as the pneumococcal vaccine at 3 and 5 months. The DTPa-IPV/Hib vaccine was also co-administered at 11 to 12 months of age.</p>
Outcomes	<p><b>Primary outcome:</b> parent-reported, physician-confirmed all-cause AOM (stratified to 1 or more AOM episodes and overall AOM)</p> <p><b>Secondary outcomes:</b> parent-reported, physician-confirmed all-cause AOM with antibiotic prescription (stratified to 1 or more AOM episodes and overall AOM), serious adverse events occurring throughout the study period</p> <p>Parents were asked by automatic text message every 2 weeks if their child had had a physician-confirmed AOM diagnosis. If no contact could be made, AOM status was checked at the next study visit.</p>
Funding sources	<p>GlaxoSmithKline Biologicals SA was the funding source and was involved in all stages of the study conduct and analysis. GlaxoSmithKline Biologicals SA also took responsibility for all costs associated with the development and publishing of this article.</p>
Declarations of interest	<p>Dr Vesikari declares that he received payment from the GlaxoSmithKline group of companies and other vaccine manufacturers for board membership, consultancy, and attending meetings; the institution of Dr Kaijalainen received grants from the GlaxoSmithKline group of companies. 7 co-authors are em-</p>



## Vesikari 2016 (Continued)

employees of the GlaxoSmithKline group of companies. Dr Hezareh is a consultant for Chiltern International for the GlaxoSmithKline group of companies. Dr Puumalainen was a GlaxoSmithKline group of companies employee during the study. 4 co-authors declare stock and stock options ownership in the GlaxoSmithKline group of companies, and 1 co-author declares shares ownership in the GlaxoSmithKline group of companies. Dr Forsten and Dr Seppä declare no conflicts of interest.

### Notes

**Participants lost to follow-up:total:** 976/5093 (19.2%) did not complete the follow-up period specified in the protocol

**Participants lost to follow-up:Tx1:** 0/1846 (0%) did not complete the follow-up period specified in the protocol

**Participants lost to follow-up:Tx2:** 371/1313 (28.3%) did not complete the follow-up period specified in the protocol

**Participants lost to follow-up:C1:** 605/1073 (56.4%) did not complete the follow-up period specified in the protocol

**Participants lost to follow-up:C2:** 0/861 (0%) did not complete the follow-up period specified in the protocol

### Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Quote: "Clusters were randomized (2:2:1:1: PHiD-CV 3+1, PHiD-CV 2+1, control 3+1, control 2+1) using a blocking scheme, stratified according to cluster size (below/above average), urbanity (urban/rural), and Tampere University Vaccine Research Centre trial enrolment"
Allocation concealment (selection bias)	Low risk	Quote: "For nested study participants, individual randomization codes were used, aligned with cluster randomization based on place of residence"
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Stated that this was a double-blind trial, but no further details provided
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Stated that this was a double-blind trial, but no further details provided
Incomplete outcome data (attrition bias) All outcomes	High risk	Substantial number of participants not included in analysis due to "randomization error"
Selective reporting (reporting bias)	Low risk	Prespecified outcomes listed in <a href="https://clinicaltrials.gov/show/NCT00839254">clinicaltrials.gov/show/NCT00839254</a> .
Other bias	Low risk	No other sources of bias identified.

Ab: antibiotics

AOM: acute otitis media

C: control

CAP: community-acquired pneumonia

CRM197-PCV7: 7-valent pneumococcal conjugate vaccine conjugated to carrier protein CRM197

CRM197-PCV9: 9-valent pneumococcal conjugate vaccine conjugated to carrier protein CRM197

DCC: daycare centre

DTaP: diphtheria-tetanus toxoid-acellular pertussis vaccine

DTP: diphtheria-tetanus toxoid-pertussis vaccine  
DTwP: diphtheria-tetanus toxoid-whole cell pertussis vaccine  
ELISA: enzyme-linked immunosorbent assay  
ENT: ear, nose, and throat  
GP: general practitioner  
HBV/placebo: hepatitis B virus vaccination plus placebo vaccine  
Hib: *Haemophilus influenzae* type b  
ICPC: International Classification of Primary Care  
IgA: immunoglobulin A  
IgG: immunoglobulin G  
IPD: invasive pneumococcal disease  
ITT: intention-to-treat  
MenC: meningococcus type C conjugate vaccine  
NaCl: sodium chloride  
OM: otitis media  
OMPC-PCV7: 7-valent pneumococcal conjugate vaccine conjugated to the outer membrane protein complex of *Neisseria meningitidis* serogroup B  
PCR: polymerase chain reaction  
PCV: pneumococcal conjugate vaccine  
PHiD-CV10: 10-valent pneumococcal conjugate vaccine conjugated to protein D (surface lipoprotein of non-typeable *Haemophilus influenzae*)  
PHiD-CV11: 11-valent pneumococcal conjugate vaccine conjugated to protein D (surface lipoprotein of non-typeable *Haemophilus influenzae*)  
PPV23: 23-valent pneumococcal polysaccharide vaccine  
RTI: respiratory tract infection  
TIV: trivalent influenza vaccine  
TIV/CRM197-PCV7: trivalent influenza vaccine plus 7-valent pneumococcal conjugate vaccine conjugated to carrier protein CRM197  
TIV/placebo: trivalent influenza vaccine plus placebo vaccine  
Tx: treatment

### Characteristics of excluded studies [ordered by study ID]

Study	Reason for exclusion
<a href="#">Gisselsson-Solen 2011</a>	No control vaccination
<a href="#">Jokinen 2012</a>	Re-analysis of the <a href="#">Eskola 2001</a> study, with no new outcome data that could be used in our review
<a href="#">Le 2007</a>	RCT studying the effect of PCV on OME
<a href="#">Palmu 2014</a>	Secondary analysis of the FinIP trial, <a href="#">Palmu 2013</a> , with no outcome data that could be used in our review
<a href="#">Palmu 2015a</a>	Re-analysis of the <a href="#">Eskola 2001</a> study, with no new outcome data that could be used in our review
<a href="#">Palmu 2015b</a>	Secondary analysis of the FinIP trial, <a href="#">Palmu 2013</a> , with no outcome data that could be used in our review
<a href="#">Palmu 2018</a>	Secondary analysis of the FinIP trial, <a href="#">Palmu 2013</a> , with no outcome data that could be used in our review
<a href="#">Roy 2011</a>	RCT studying the effect of PCV on suppurative otitis media (abstract of conference meeting)
<a href="#">Sarasoja 2013</a>	Re-analysis of the <a href="#">Eskola 2001</a> study, with no new outcome data that could be used in our review

OME: otitis media with effusion  
PCV: pneumococcal conjugate vaccine  
RCT: randomised controlled trial

## ADDITIONAL TABLES

**Table 1. Effect of pneumococcal conjugate vaccination on frequency of all-cause acute otitis media episodes**

	Intention-to-treat				Per-protocol			
	Episodes/person-year		Incidence rate difference - episodes per person-year (95% CI)	VE expressed as relative reduction in risk (95% CI) <sup>a</sup>	Episodes/person-year		Incidence rate difference - episodes per person-year (95% CI)	VE expressed as relative reduction in risk (95% CI) <sup>a</sup>
	Treatment	Control			Treatment	Control		
PCV administered in early infancy								
CRM197-PCV7								
Black 2000	-	-	-	6% (4% to 9%)	-	-	-	7% (4% to 10%)
Fireman 2003	-	-	-	6% (4% to 8%)	-	-	-	7% (4% to 9%)
Eskola 2001	-	-	-	-	1.16	1.24	-0.08 <sup>d</sup>	6% (-4% to 16%)
O'Brien 2008 <sup>b</sup>	1.43	1.36	0.07 (-0.05 to 0.18)	-5% (-25% to 12%) <sup>c</sup>	1.35	1.35	0.00 (-0.13 to 0.14)	0% (-21% to 17%)
OMPC-PCV7								
Kilpi 2003	-	-	-	-	-	-	-	-1% <sup>h</sup> (-12% to 10%)
PHiD-PC10 and PHiD-PC11								
Tregnaghi 2014	0.03	0.04	-0.01 (-0.01 to 0.00)	15% (-1% to 28%)	-	-	-	13% (-5% to 28%)
Sáez-Llorens 2017								
Vesikari 2016 <sup>b</sup>	-	-	-	-	0.99	1.01	-0.02 <sup>d</sup>	6% (-6% to 17%)
Karppinen 2019 <sup>e</sup>	-	-	-	-	1.0	1.3	-0.3 (-0.7 to 0.1)	23% (0% to 40%)
Prymula 2006	-	-	-	-	0.08	0.13	-0.04 <sup>d</sup>	34% (21% to 44%)

**Table 1. Effect of pneumococcal conjugate vaccination on frequency of all-cause acute otitis media episodes** (Continued)

**PCV administered at a later age**

**CRM197-PCV7 followed by PPV23**

Veenhoven 2003	-	-	-	-25% (-57% to 1%)	1.1	0.83	-0.27 <sup>d</sup>	-29% <sup>h</sup> (-62% to -2%)
van Kempen 2006	-	-	-	-	0.78	0.67	-0.11 <sup>d</sup>	-16% <sup>h</sup> (-96% to 31%)

**CRM197-PCV7/TIV**

Jansen 2008	-	-	-	-	-	-	-	57% (6% to 80%) <sup>f</sup>
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**CRM197-PCV9**

Dagan 2001	-	-	-	-	0.66	0.79	-0.14 (-0.29 to 0.02)	17% (-2% to 33%)
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CI: confidence interval

CRM197-PCV7: 7-valent pneumococcal conjugate vaccine conjugated to carrier protein CRM197

CRM197-PCV7/TIV: trivalent influenza vaccine plus 7-valent pneumococcal conjugate vaccine conjugated to carrier protein CRM197

CRM197-PCV9: 9-valent pneumococcal conjugate vaccine conjugated to carrier protein CRM197

HBV: hepatitis B virus

OMPC-PCV7: 7-valent pneumococcal conjugate vaccine conjugated to the outer membrane protein complex of *Neisseria meningitidis* serogroup B

PCV: pneumococcal conjugate vaccine

PHiD-CV10: 10-valent pneumococcal conjugate vaccine conjugated to protein D (surface lipoprotein of non-typeable *Haemophilus influenzae*)

PHiD-CV11: 11-valent pneumococcal conjugate vaccine conjugated to protein D (surface lipoprotein of non-typeable *Haemophilus influenzae*)

PPV23: 23-valent pneumococcal polysaccharide vaccine

TIV: trivalent influenza vaccine

VE: vaccine efficacy

<sup>a</sup>Positive effect estimates indicate a relative reduction in the risk (e.g. 6% means that the vaccine *reduces* the risk by 6%); negative effect estimates indicate a relative increase in the risk (e.g. -5% means that the vaccine *increases* the risk by 5%).

<sup>b</sup>Cluster-randomised controlled trial.

<sup>c</sup>Defined as primary efficacy analysis. Analysis is not entirely according to intention-to-treat principle, as 88/944 children were not included in the analysis due to not meeting strict chart review criteria.

<sup>d</sup>95% CI could not be calculated, as person-time across treatment groups was not reported.

<sup>e</sup>Respiratory tract infections with acute otitis media was used as the outcome measure. The PHiD-CV10 and control vaccine groups were statistically different from each other in terms of type of residential area, presence of older siblings, and socioeconomic status of the family.

<sup>f</sup>Index group: CRM197-PCV7/TIV, control: HBV/placebo; VE placebo/TIV versus HBV/placebo: 71% (95% CI 30% to 88%), that is larger VE placebo/TIV versus HBV/placebo than CRM197-PCV7/TIV versus HBV/placebo.

<sup>h</sup>negative values for VE expressed as relative reduction in risk represent an increase in the risk for acute otitis media.

**Table 2. Effect of pneumococcal conjugate vaccination on frequency of pneumococcal acute otitis media episodes**

	Intention-to-treat				Per-protocol			
	VE expressed as relative reduction in risk (95% CI)				VE expressed as relative reduction in risk (95% CI)			
	Pneumo- coccal AOM	Vaccine-type AOM	Cross-reac- tive-type AOM	Non-vac- cine-type AOM	Pneumococcal AOM	Vaccine-type AOM	Cross-reac- tive-type AOM	Non-vac- cine-type AOM
<b>PCV administered in infancy</b>								
<b>CRM197-PCV7</b>								
Black 2000 <sup>a</sup>	-	65% P = 0.04	-	-	-	67% P = 0.08	-	-
Fireman 2003	-	-	-	-	-	-	-	-
Eskola 2001	-	54% (41% to 64%)	-	-	34% (21% to 45%)	57% (44% to 67%)	51% (27% to 67%)	-33% <sup>d</sup> (-80% to 1%)
Palmu 2009 <sup>b</sup>	-	-	-	-	20% (7% to 31%)	-	-	-
O'Brien 2008 <sup>a,c</sup>	-	64% (-34% to 90%)	-	-	-	-	-	-
<b>OMPC-PCV7</b>								
Kilpi 2003	-	-	-	-	25% (11% to 37%)	56% (44% to 66%)	-5% <sup>d</sup> (-47% to 25%)	-27% <sup>d</sup> (-70% to 6%)
<b>PHiD-PC10 and PHiD-PC11</b>								
Tregnaghi 2014	53% (16% to 74%)	70% (30% to 87%)	29% (-123% to 77%)	15% (-153% to 71%)	56% (13% to 78%)	67% (17% to 87%)	26% (-232% to 83%)	26% (-231% to 83%)
Sáez-Llorens 2017								
Vesikari 2016 <sup>c</sup>	-	-	-	-	-	-	-	-
Prymula 2006	-	-	-	-	52% (37% to 63%)	58% (41% to 69%)	66% (22% to 85%)	9% (-64% to 49%)
<b>PCV administered at a later age</b>								



**Table 2. Effect of pneumococcal conjugate vaccination on frequency of pneumococcal acute otitis media episodes** (Continued)

**CRM197-PCV7 followed by PPV23**

Veenhoven 2003	-	-	-	-	34% P = 0.22	52% P = 0.21	-	21% P = 0.44
van Kempen 2006	-	-	-	-	-	-	-	-

**CRM197-PCV7/TIV**

Jansen 2008	-	-	-	-	-	-	-	-
-------------	---	---	---	---	---	---	---	---

**CRM197-PCV9**

Dagan 2001	-	-	-	-	-	-	-	-
------------	---	---	---	---	---	---	---	---

AOM: acute otitis media

CI: confidence interval

CRM197-PCV7: 7-valent pneumococcal conjugate vaccine conjugated to carrier protein CRM197

CRM197-PCV7/TIV: trivalent influenza vaccine plus 7-valent pneumococcal conjugate vaccine conjugated to carrier protein CRM197

CRM197-PCV9: 9-valent pneumococcal conjugate vaccine conjugated to carrier protein CRM197

OMPC-PCV7: 7-valent pneumococcal conjugate vaccine conjugated to the outer membrane protein complex of *Neisseria meningitidis* serogroup B

PCV: pneumococcal conjugate vaccine

PPV23: 23-valent pneumococcal polysaccharide vaccine

TIV: trivalent influenza vaccine

VE: vaccine efficacy

<sup>a</sup>Middle ear fluid collected from spontaneous draining ears; in the other studies middle ear fluid was routinely collected during AOM episodes through paracentesis.

<sup>b</sup>Additional analysis of [Eskola 2001](#) including pneumococcal AOM by a positive culture or polymerase chain reaction (PCR).

<sup>c</sup>Cluster-randomised controlled trial.

<sup>d</sup>negative values represent an increase in the risk of AOM.

**Table 3. Effect of pneumococcal conjugate vaccination on frequency of recurrent acute otitis media**

	Intention-to-treat	Per-protocol
	VE expressed as relative reduction in risk (95% CI)	VE expressed as relative reduction in risk (95% CI)
<b>PCV administered in infancy</b>		
<b>CRM197-PCV7</b>		
Black 2000	9% (4% to 14%)	9% (3% to 15%)
Fireman 2003	10% (7% to 13%)	-
Eskola 2001	9% (-12% to 27%)	16% (-6% to 35%)
O'Brien 2008 <sup>a</sup>	-	-
<b>OMPC-PCV7</b>		
Kilpi 2003	-	-
<b>PHiD-PC10 and PHiD-PC11</b>		
Tregnaghi 2014	-	-
Sáez-Llorens 2017	-	-
Vesikari 2016 <sup>a</sup>	-	-
Prymula 2006	-	56% (-2% to 81%)
<b>PCV administered at a later age</b>		
<b>CRM197-PCV7 followed by PPV23</b>		
Veenhoven 2003	-	-
van Kempen 2006	-	-
<b>CRM197-PCV7/TIV</b>		
Jansen 2008	-	-
<b>CRM197-PCV9</b>		
Dagan 2001	-	-

CI: confidence interval

CRM197-PCV7: 7-valent pneumococcal conjugate vaccine conjugated to carrier protein CRM197

CRM197-PCV7/TIV: trivalent influenza vaccine plus 7-valent pneumococcal conjugate vaccine conjugated to carrier protein CRM197

CRM197-PCV9: 9-valent pneumococcal conjugate vaccine conjugated to carrier protein CRM197

OMPC-PCV7: 7-valent pneumococcal conjugate vaccine conjugated to the outer membrane protein complex of *Neisseria meningitidis* serogroup B

PCV: pneumococcal conjugate vaccine

PPV23: 23-valent pneumococcal polysaccharide vaccine

TIV: trivalent influenza vaccine

VE: vaccine efficacy

<sup>a</sup>Cluster-randomised controlled trial

**Table 4. Adverse effects**

Study ID	No. of participants	PCV type	Redness	Swelling	Pain/tenderness	Fever	Serious adverse events
Black 2000/ Fireman 2003	37,868	CRM197-PCV7	<p>Depending on timing of dose, redness occurred in around 10% to 14% of children receiving CRM197-PCV7 versus 5% to 9% of children receiving MenC vaccination.</p> <p>More severe redness (&gt; 3 cm) occurred in 0% to 0.6% of children receiving CRM197-PCV7, and did not differ significantly between CRM197-PCV7 and MenC groups.</p>	<p>Depending on timing of dose, swelling occurred in around 10% to 12% of children receiving CRM197-PCV7 versus 3% to 8% of children receiving MenC vaccination.</p> <p>More severe swelling (&gt; 3 cm) occurred in 0.1% to 0.6% of children receiving CRM197-PCV7, and did not differ significantly between CRM197-PCV7 and MenC groups.</p>	<p>Depending on timing of dose, tenderness was reported in 15% to 23% of children receiving CRM197-PCV7, and did not differ significantly between CRM197-PCV7 and MenC groups.</p>	<p>Depending on timing of dose, fever &gt; 38 °C occurred in around 15% to 24% of children receiving CRM197-PCV7 versus 9% to 17% of children receiving MenC vaccination.</p> <p>Fever (&gt; 39 °C) occurred in 0.9% to 2.5% of children receiving</p>	<p>No severe adverse events related to vaccination resulting in hospitalisation, emergency, or clinic visits were reported.</p>

**Table 4. Adverse effects** (Continued)

							CRM197-PCV7, and did not differ significantly between CRM197-PCV7 and MenC groups.
Dagan 2001	264	CRM197-PCV9	Depending on timing of dose, redness occurred in 5% to 6% of children receiving CRM197-PCV9 versus 0% to 5% of children receiving MenC vaccination.	Depending on timing of dose, swelling occurred in 7% to 12% of children receiving CRM197-PCV9 versus 0% to 5% of children receiving MenC vaccination.	Depending on timing of dose, tenderness was reported in 25% to 38% of children receiving CRM197-PCV9 versus 0% to 8% of children receiving MenC vaccination.	Depending on timing of dose, fever > 38 °C occurred in around 15% to 44% of children receiving CRM197-PCV9 versus 8% to 25% of children receiving MenC vaccination.	Not reported

**Table 4. Adverse effects** (Continued)

						Fever (> 39.5 °C) oc- curred in only 1 child receiv- ing CRM197- PCV9 versus 3 chil- dren receiv- ing MenC vacci- nation.	
Eskola 2001/ Palmu 2009	1662	CRM197-PCV7	Depend- ing on tim- ing of dose, redness oc- curred in 14% to 20% of children receiving CRM197- PCV7 ver- sus 9% to 16% of chil- dren receiv- ing hepatitis vaccines.	Depend- ing on tim- ing of dose, swelling oc- curred in 5% to 6% of children receiving CRM197- PCV7 versus 2% to 6% of children re- ceiving he- patitis vac- cines.	Depending on timing of dose, pain was report- ed in 3% to 8% of chil- dren receiving CRM197-PCV7 versus 2% to 3% of children receiving he- patitis vac- cines.	Fever (> 39 °C) oc- curred in 0.4% to 2.0% of chil- dren receiv- ing CRM197- PCV7 versus 0.2% to 1.7% of chil- dren receiv- ing he- pati- tis vac- cines.	No significant differences between vaccine groups were ob- served for unexpected events (6 versus 4 events).  1 child in the CRM197-PCV7 group died from bowel obstruc- tion, necrosis, and shock at the age of 8 months (85 days af- ter administration of third dose), but death was assessed as unrelated to study vaccine (autopsy revealed mesenteric defects with volvulus and other congenital abnormalities).



**Table 4. Adverse effects** (Continued)

			nificantly between CRM197- PCV7 and hepatitis vaccine groups.	nificantly between CRM197- PCV7 and hepatitis vaccine groups.			
Jansen 2008	579	CRM197- PCV7/TIV	-	-	-	-	Quote: "In general, the vaccinations were well-tolerated, and no immediate or severe adverse events were recorded."
Kilpi 2003	1666	OMPC-PCV7	OMPC-PCV7 caused local reactions within 3 days of each dose more often than the hepB vaccine (data not shown).	OMPC-PCV7 caused local reactions within 3 days of each dose more often than the hepB vaccine (data not shown).	Not reported	Not reported	There were no statistically significant differences in the occurrence of any diagnosis among individuals who experienced serious adverse events between the 2 vaccine groups.  1 child in the OMPC-PCV7 group died from volvulus due to bowel obstruction. Death was assessed as unrelated to study vaccine.
Prymula 2006	4968	PHiD-CV11	Not reported	Not reported	Not reported	Not reported	The percentages of infants with unsolicited symptoms that were judged to be causally related to vaccination were similar in the PHiD-CV11 and hepA groups (2.5% versus 3.0%). 14 serious adverse events were judged to be causally related to vaccination: 8 occurred in children receiving PHiD-CV11 vaccination (7 after co-administration with Infanrix hexa and 1 after PHiD-CV11 booster) versus 6 in children receiving hepatitis A control vaccine (7 after co-administration with Infanrix hexa and 1 after hepatitis A booster with Infanrix hexa). All events, apart from 1 case of epilepsy in the hepatitis A group, resolved without sequelae.  4 children died during the study, 1 in the PHiD-CV11 group (8 months after third dose, diagnosis of epilepsy was made; 25 months after the third dose the child had grand mal epilepsy and died from suffocation). None of the deaths were regarded by the investigators as related to the study vaccine.

**Table 4. Adverse effects** (Continued)

Tregnaghi 2014/Sáez- Llorens 2017	23,821	PHiD-CV10	Not report- ed	Not report- ed	Not reported	Not re- ported	Serious adverse events did not differ significantly between PHiD-CV10 and hepatitis control vaccines (21.5% versus 22.6%). Only 1 event (in the control group) was judged to be causally related to vaccination by the investigator, and it resolved without sequelae.  19 children died in the PHiD-CV10 group (0.16%) versus 26 in the control group (0.22%). None of the deaths were considered by the investigator to be causally related to vaccination.
Veenhoven 2003	383	CRM197-PCV7	Not report- ed	Not report- ed	Not reported	Not re- ported	No serious adverse events were noted after administration of CRM197-PCV7 or hepatitis control vaccines.
Vesikari 2016	6178	PHiD-CV10	Not report- ed	Not report- ed	Not reported	Not re- ported	Serious adverse events considered by the investigator to be causally related to vaccination were reported in 4 infants in the PHiD-CV10 group (all in 3 + 1 group: sepsis with non-specified aetiology in 1 infant, pyrexia in 1 infant, convulsion in 2 infants) and in 2 infants in hepB group (petit mal epilepsy in 1 infant and pyrexia in 1 infant).  1 fatal serious adverse event (sudden infant death, not considered to be vaccination related) was reported in the PHiD-CV10 (2 + 1) group.

CRM197-PCV7: 7-valent pneumococcal conjugate vaccine conjugated to carrier protein CRM197

CRM197-PCV7/TIV: trivalent influenza vaccine plus 7-valent pneumococcal conjugate vaccine conjugated to carrier protein CRM197

CRM197-PCV9: 9-valent pneumococcal conjugate vaccine conjugated to carrier protein CRM197

hepA: hepatitis A

hepB: hepatitis B

MenC: meningococcus type C

OMPC-PCV7: 7-valent pneumococcal conjugate vaccine conjugated to the outer membrane protein complex of *Neisseria meningitidis* serogroup B

PHiD-CV10: 10-valent pneumococcal conjugate vaccine conjugated to protein D (surface lipoprotein of non-typeable *Haemophilus influenzae*)

PHiD-CV11: 11-valent pneumococcal conjugate vaccine conjugated to protein D (surface lipoprotein of non-typeable *Haemophilus influenzae*)

TIV: trivalent influenza vaccine

## APPENDICES

### Appendix 1. Previous search details

For the 2014 version of the review we searched the Cochrane Central Register of Controlled Trials (CENTRAL) (2013, Issue 11), which contains the Cochrane Acute Respiratory Infections Group's Specialised Register; MEDLINE (1995 to November week 3, 2013); Embase (1995 to December 2013); CINAHL (2007 to December 2013); LILACS (2007 to December 2013) and Web of Science (2007 to December 2013).

We used the following search strategy to search CENTRAL and MEDLINE. We combined the MEDLINE search with the Cochrane Highly Sensitive Search Strategy for identifying randomised trials in MEDLINE: sensitivity- and precision-maximising version (2008 revision); Ovid format (Lefebvre 2011). We adapted the search strategy to search Embase, CINAHL, LILACS and Web of Science.

#### MEDLINE (Ovid)

```
1 exp Otitis Media/
2 otitis media.tw.
3 aom.tw.
4 or/1-3
5 Pneumococcal Vaccines/
6 Vaccines, Conjugate/
7 Bacterial Vaccines/
8 (pneumococc* adj5 (vaccin* or conjugat* or immuni*)).tw,nm.
9 pcv*.tw,nm.
10 or/5-9
11 4 and 10
```

For the 2009 version of the review we searched the Cochrane Central Register of Controlled Trials (CENTRAL) (*The Cochrane Library* 2007, issue 2), which contains the Cochrane Acute Respiratory Infections Group's Specialised Register; MEDLINE (January 1995 to November 2007); and EMBASE (January 1995 to November 2007).

We used the following search strategy for searching MEDLINE and CENTRAL and modified terms for searching EMBASE.

#### MEDLINE

```
#1 explode 'bacterial-vaccine' / all subheadings
#2 explode 'bacterial AND vaccine' / all subheadings
#3 explode 'Pneumococcus-vaccine' / all subheadings
#4 pneumococc* near immunity*
#5 pneumococc* near vaccin*
#6 #1 or #2 or #3 or #4 or #5
#7 explode 'otitis media' / all subheadings
#8 (otitis media in ti) or (otitis media in ab)
#9 #7 or #8
#10 #6 or #9
#11 explode 'randomised-controlled-trial' / all subheadings
#12 explode 'controlled-study' / all subheadings
#13 explode 'randomisation' / all subheadings
#14 explode 'single-blind-procedure' / all subheadings
#15 explode 'double-blind-procedure' / all subheadings
#16 explode 'crossover-procedure' / all subheadings
#17 explode 'phase-3-clinical-trial' / all subheadings
#18 (control* near trial*) in ti
#19 (control* near trial*) in ab
#20 (singl* or doubl* or trebl* or tripl*) near ((blind* or mask*) in ti)
#21 (singl* or doubl* or trebl* or tripl*) near ((blind* or mask*) in ab)
#22 random* near ((allocat* or assign*) in ti)
#23 random* near ((alsocat* or assign*) in ab)
#24 #11 or #12 or #13 or #14 or #15 or #16 or #17 or #18 or #19 or #20 or #21 or #22 or #23
#25 #10 and #24
```

### Appendix 2. MEDLINE (Ovid) search strategy

```
1 exp Otitis Media/
2 otitis media.tw.
```

3 (aom or om or ome or csom).tw.  
4 glue ear.tw.  
5 (otorrhea\* or otorrhea\*).tw.  
6 (middle ear adj5 (infect\* or inflam\* or effus\*)).tw.  
7 or/1-6  
8 exp Pneumococcal Vaccines/  
9 Vaccines, Conjugate/  
10 Bacterial Vaccines/  
11 (pneumococc\* adj5 (vaccin\* or conjugat\* or immuni\*)).tw,nm.  
12 (pcv\* or 5PPV\* or 7PCV\* or 9PCV\* or 10PCV\* or 11PCV\* or 13PCV\* or 15PCV\* or synflorix).tw,nm.  
13 or/8-12  
14 7 and 13

### Appendix 3. Embase (Elsevier) search strategy

#17. #12 AND #16  
#16. #13 OR #14 OR #15  
#15. random\*:ab,ti OR placebo\*:ab,ti OR factorial\*:ab,ti OR crossover\*:ab,ti OR 'cross over':ab,ti OR 'cross-over':ab,ti OR volunteer\*:ab,ti  
OR assign\*:ab,ti OR allocat\*:ab,ti OR ((singl\* OR doubl\*) NEAR/1 blind\*):ab,ti  
#14. 'single blind procedure'/exp OR 'double blind procedure'/exp OR 'crossover procedure'/exp  
#13. 'randomized controlled trial'/exp  
#12. #6 AND #11  
#11. #7 OR #8 OR #9 OR #10  
#10. (pcv\* or 5PPV\* or 7PCV\* or 9PCV\* or 10PCV\* or 11PCV\* or 13PCV\* or 15PCV\* or synflorix):ab,ti,tn  
#9. (pneumococc\* NEAR/5 (vaccin\* OR conjugat\* OR immuni\*)):ab,ti,tn  
#8. 'bacterial vaccine'/de  
#7. 'pneumococcus vaccine'/de  
#6. #1 OR #2 OR #3 OR #4 OR #5  
#5. ("middle ear" NEAR/5 (infect\* OR inflam\* OR effus\*)):ab,ti  
#4. otorrhea\*:ab,ti OR otorrhea\*:ab,ti  
#3. 'glue ear':ab,ti  
#2. 'otitis media':ab,ti OR aom:ab,ti OR om:ab,ti OR ome:ab,ti OR csom:ab,ti  
#1. 'otitis media'/exp

### Appendix 4. CINAHL (EBSCO) search strategy

S24 S14 and S23  
S23 S15 or S16 or S17 or S18 or S19 or S20 or S21 or S22  
S22 (MH "Quantitative Studies")  
S21 (MH "Placebos")  
S20 TI placebo\* OR AB placebo\*  
S19 TI random\* OR AB random\*  
S18 TI ( singl\* blind\* or doubl\* blind\* or tripl\* blind\* or trebl\* blind\* or singl\* mask\* or doubl\* mask\* or tripl\* mask\* or trebl\* mask\* ) OR  
AB (singl\* blind\* or doubl\* blind\* or tripl\* blind\* or trebl\* blind\* or singl\* mask\* or doubl\* mask\* or tripl\* mask\* or trebl\* mask\* )  
S17 TI clinic\* trial\* OR AB clinic\* trial\*  
S16 PT clinical trial  
S15 (MH "Clinical Trials+")  
S14 S6 and S13  
S13 S7 or S8 or S9 or S10 or S11 or S12  
S12 (TI pcv\* OR AB pcv\* OR TI 5PPV\* OR AB 5PPV\* OR TI 7PCV\* OR AB 7PCV\* OR TI 9PCV\* OR AB 9PCV\* OR TI 10PCV\* OR AB 10PCV\* OR TI  
11PCV\* OR AB 11PCV\* OR TI 13PCV\* OR AB 13PCV\* OR TI 15PCV\* OR AB 15PCV\* OR TI synflorix OR AB synflorix)  
S11 TI pneumococc\* N5 immuni\* OR AB pneumococc\* N5 immuni\*  
S10 TI pneumococc\* N5 conjugat\* OR AB pneumococc\* N5 conjugat\*  
S9 TI pneumococc\* N5 vaccin\* OR AB pneumococc\* N5 vaccin\*  
S8 (MH "Bacterial Vaccines")  
S7 (MH "Pneumococcal Vaccine")  
S6 S1 or S2 or S3 or S4 or S5  
S5 ((TI "middle ear" OR AB "middle ear") N5 (TI infect\* OR AB infect\* OR TI inflam\* OR AB inflam\* OR TI effus\* OR AB effus\*))  
S4 TI (otorrhea\* or otorrhea\*) OR AB (otorrhea\* or otorrhea\*)  
S3 TI glue ear OR AB glue ear  
S2 TI ( otitis media or aom or ome or csom) OR AB ( otitis media or aom or ome or csom)  
S1 (MH "Otitis Media+")

## Appendix 5. LILACS (BIREME) search strategy

```
tw:((mh:"otitis media" OR "otitis media" OR "Otitis Média" OR mh:c09.218.705.663* OR aom OR om OR ome OR csom OR "glue ear" OR otorrhea* OR otorrhoea* OR (middle ear AND (infect* OR inflam* OR effus*))) AND (mh:"Pneumococcal Vaccines" OR "Vacunas Neumococicas" OR "Vacinas Pneumocócicas" OR mh:"Vaccines, Conjugate" OR "Vacunas Conjugadas" OR "Vacinas Conjugadas" OR mh:"Bacterial Vaccines" OR "Vacunas Bacterianas" OR "Vacinas Bacterianas" OR "pneumococcal vaccine" OR "pneumococcal vaccines" OR "conjugate vaccines" OR "conjugate vaccine" OR pcv* OR 5ppv* OR 7pcv* OR 9pcv* OR 10pcv* OR 11pcv* OR 13pcv* OR 15pcv* OR synflorix)) AND (instance:"regional") AND (db:"LILACS"))
```

## Appendix 6. Web of Science (Clarivate Analytics) search strategy

# 5	113	#4 AND #3  Indexes=SCI-EXPANDED, CPCI-S, CCR-EXPANDED Timespan=2013-2018
# 4	719,817	<b>TOPIC:</b> ((random* or placebo* or allocat* or crossover* or "cross over" or ((doubl* or singl*) NEAR/1 (blind* or mask*)))) OR <b>TITLE:</b> (trial)  Indexes=SCI-EXPANDED, CPCI-S, CCR-EXPANDED Timespan=2013-2018
# 3	498	#2 AND #1  Indexes=SCI-EXPANDED, CPCI-S, CCR-EXPANDED Timespan=2013-2018
# 2	6,232	<b>TOPIC:</b> ((pneumococc* NEAR/5 (vaccin* or conjugat* or immuni*))) OR <b>TOPIC:</b> ((pcv* or 5PPV* or 7PCV* or 9PCV* or 10PCV* or 11PCV* or 13PCV* or 15PCV* or synflorix))  Indexes=SCI-EXPANDED, CPCI-S, CCR-EXPANDED Timespan=2013-2018
# 1	15,363	<b>TOPIC:</b> ("otitis media" or aom or om or ome or csom or "glue ear" or otorrhea* or otorrhoea* or ("middle ear" NEAR/5 (infect* or inflam* or effus*)))  Indexes=SCI-EXPANDED, CPCI-S, CCR-EXPANDED Timespan=2013-2018

## Appendix 7. ClinicalTrials.gov search strategy

("otitis media" OR aom OR om OR ome OR csom OR "glue ear" OR otorrhea OR otorrhoea OR "middle ear infection" OR "middle ear infections") AND (pneumococcal OR pneumococcus) AND (vaccine OR vaccines OR vaccinating OR vaccination OR vaccinations)

("otitis media" OR aom OR om OR ome OR csom OR "glue ear" OR otorrhea OR otorrhoea OR "middle ear infection" OR "middle ear infections") AND (pneumococcal OR pneumococcus) AND (conjugate OR conjugated OR immunisation)

("otitis media" OR aom OR om OR ome OR csom OR "glue ear" OR otorrhea OR otorrhoea OR "middle ear infection" OR "middle ear infections") AND (pneumococcal OR pneumococcus) AND (immunisations OR immunised OR immunising)

(Search has been split into 3 to work better on the ClinicalTrials.gov search portal)

## Appendix 8. WHO ICTRP search strategy

pneumococc\* AND vaccin\* AND otitis media OR pneumococc\* AND conjugat\* AND otitis media OR pneumococc\* AND immuni\* AND otitis media OR pcv\* AND otitis media OR pneumococc\* AND vaccin\* AND aom OR pneumococc\* AND conjugat\* AND aom OR pneumococc\*

AND immuni\* AND aom OR pcv\* AND aom OR pneumococc\* AND vaccin\* AND middle ear infection\* OR pneumococc\* AND conjugat\* AND middle ear infection\* OR pneumococc\* AND immuni\* AND middle ear infection\* OR pcv\* AND middle ear infection\*

## WHAT'S NEW

Date	Event	Description
11 June 2020	New citation required but conclusions have not changed	The included trial publication (secondary analysis of a previously included trial) did not change the conclusions.
11 June 2020	New search has been performed	<p>Two new review authors joined the team. One new review author (VL) conducted systematic searches for randomised controlled trials and controlled clinical trials to update the review. The other new review author (JHLdS) independently assessed risk of bias, determined the certainty of the evidence using the GRADE framework, and performed data extraction of all eligible studies. Any discrepancies in results for this update were discussed with a second review author (RPV) and where needed with a third review author (RAMJD).</p> <p>We changed some of the GRADE assessments in this review update. On reflection, we felt that our GRADE assessments were incorrect in the last publication; our confidence in the effect estimates for some of the outcomes was too optimistic in previous versions.</p> <p>We updated the searches (March 2019 to June 2020) and included one new trial publication (Karppinen 2019) as a secondary analysis of <a href="#">Vesikari 2016</a> (FinIP vaccine trial). No further studies were excluded.</p> <p>We did not identify any ongoing studies.</p>

## HISTORY

Protocol first published: Issue 2, 1999

Review first published: Issue 2, 2002

Date	Event	Description
9 March 2020	Amended	The authors' <a href="#">Declarations of interest</a> have been updated to reflect the review's compliance with the <a href="#">Cochrane conflict of interest policy</a> , which includes the relevant parts of the <a href="#">Cochrane Commercial Sponsorship Policy</a> .
29 March 2019	New search has been performed	<p>We updated the searches and included three new publications: two new randomised controlled trials (RCTs) (<a href="#">Tregnaghi 2014</a>; <a href="#">Vesikari 2016</a>), and one new trial publication, Sáez-Llorens 2017, that reported outcome data relevant for this review as part of a secondary analysis of <a href="#">Tregnaghi 2014</a>.</p> <p>We excluded five new trial publications (secondary analyses of previously included RCTs) (<a href="#">Palmu 2014</a>; <a href="#">Palmu 2015a</a>; <a href="#">Palmu 2015b</a>; <a href="#">Palmu 2018</a>; <a href="#">Sarasoja 2013</a>), since they did not report outcome data relevant to this review.</p> <p>We did not identify any ongoing studies.</p>



Date	Event	Description
		<p>The 14 publications included in this review originate from 11 RCTs (60,733 children) in total: <a href="#">Black 2000</a>/<a href="#">Fireman 2003</a>; <a href="#">Dagan 2001</a>; <a href="#">Eskola 2001</a>/<a href="#">Palmu 2009</a>; <a href="#">Jansen 2008</a>; <a href="#">Kilpi 2003</a>; <a href="#">O'Brien 2008</a>; <a href="#">Prymula 2006</a>; <a href="#">Tregnaghi 2014</a>/<a href="#">Sáez-Llorens 2017</a>; <a href="#">van Kempen 2006</a>; <a href="#">Veenhoven 2003</a>; <a href="#">Vesikari 2016</a>.</p> <p>Seven trials (59,415 children) included infants who predominantly received primary vaccinations before six months of age (<a href="#">Black 2000</a>/<a href="#">Fireman 2003</a>; <a href="#">Eskola 2001</a>/<a href="#">Palmu 2009</a>; <a href="#">Kilpi 2003</a>; <a href="#">O'Brien 2008</a>; <a href="#">Prymula 2006</a>; <a href="#">Tregnaghi 2014</a>/<a href="#">Sáez-Llorens 2017</a>; <a href="#">Vesikari 2016</a>), whilst the other four trials (1318 children) assessed the effects of PCVs administered at a later age on AOM in either healthy infants, <a href="#">Dagan 2001</a>, or in children with a history of respiratory illness or frequent AOM (<a href="#">Jansen 2008</a>; <a href="#">van Kempen 2006</a>; <a href="#">Veenhoven 2003</a>).</p> <p>We added 'adverse effects' as a co-primary outcome.</p>
29 March 2019	New citation required and conclusions have changed	<p>With this 2019 update, further information on the effect of PHiD-CV10/11 for the prevention of acute otitis media (AOM) has become available.</p> <p>Administration of the licenced CRM197-PCV7 and PHiD-CV10 during early infancy is associated with large relative risk reductions in pneumococcal AOM. However, the effects of these vaccines on all-cause AOM is far more uncertain. We found no evidence of a beneficial effect on all-cause AOM of administering pneumococcal conjugate vaccines (PCVs) in high-risk infants, after early infancy (children aged one year and above), and in older children with a history of respiratory illness. Compared to control vaccines, PCVs were associated with an increase in mild local reactions (redness, swelling), fever, and pain/tenderness, but we found no evidence of a difference in more severe local reactions, fever, or serious adverse events judged to be causally related to vaccination.</p>
21 February 2014	New search has been performed	<p>With this update, more precise information on the effect of PCV7 for the prevention of otitis media has become available. We judged the quality of the evidence for PCV7 in both early infancy and older children to be high, with further research very unlikely to change our confidence in the estimate of effect.</p> <p>Based on current evidence of the effects of PCVs for preventing AOM, the licenced 7-valent PCV has modest beneficial effects in healthy infants with a low baseline risk of AOM. Administering PCV7 in high-risk infants, after early infancy, and in older children with a history of AOM appears to have no benefit in preventing further episodes.</p> <p>Several RCTs with different (newly licenced, multivalent) PCVs administered during early infancy to establish their effects on AOM are currently ongoing. The results of these studies may provide a better understanding of the role of the newly licenced, multivalent PCVs in preventing AOM. Also, the impact of the carrier protein D, as used in certain pneumococcal vaccines for AOM, needs to be further established.</p>
3 December 2013	New search has been performed	Three new review authors joined the team to update the review.

Date	Event	Description
		<p>The updated search (November 2007 to December 2013) retrieved 171 records. After removal of duplicates, 165 records remained. After full-text review, three new publications were included in the review (Palmu 2009; Prymula 2006; van Kempen 2006). One study was an additional analysis of the previous included Eskola 2001 study.</p> <p>We identified five ongoing RCTs (NCT00466947; NCT00861380; NCT01545375; NCT01735084; NCT01174849).</p> <p>The 11 studies included in this review concerned a total of nine RCTs: (1) Black 2000/Fireman 2003; (2) Dagan 2001; (3) Eskola 2001/Palmu 2009; (4) Kilpi 2003; (5) Prymula 2006; (6) van Kempen 2006; (7) Veenhoven 2003; (8) Jansen 2008; and (9) O'Brien 2008. Five trials (n = 47,108) included healthy infants and studied the effect of PCV administered in early infancy on otitis media (OM) (Black 2000/Fireman 2003; Eskola 2001/Palmu 2009; Kilpi 2003; O'Brien 2008; Prymula 2006), whilst the other four trials (n = 1318) assessed the effects of PCV administered at a later age on OM in either healthy infants, Dagan 2001, or in children with a known history of respiratory disease including OM (Jansen 2008; van Kempen 2006; Veenhoven 2003).</p>
29 April 2008	New citation required but conclusions have not changed	New review authors
28 April 2008	Amended	Converted to new review format
15 November 2007	New search has been performed	Searches conducted.
26 November 2003	New citation required and conclusions have changed	Substantive amendment
29 June 2003	New search has been performed	Searches conducted.
19 August 2000	New search has been performed	Searches conducted.

## CONTRIBUTIONS OF AUTHORS

For the 2019 update, Alexandre C Fortanier and Roderick P Venekamp co-ordinated the review, were involved in data collection, and performed the 'Risk of bias' assessment and analysis and interpretation of the data. For the 2020 update, two new review authors joined the team. One of these review authors, Vittoria Lutje, conducted systematic searches for randomised controlled trials and controlled clinical trials. The other new review author, Joline LH de Sévaux, independently reviewed 'Risk of bias' and GRADE assessments and data extraction of all eligible studies. Any discrepancies with the results described in the 2019 update were discussed with Roderick P Venekamp, and where needed with Roger AMJ Damoiseaux. All review authors reviewed the manuscript and approved the final version of the review.

## DECLARATIONS OF INTEREST

Joline LH de Sévaux: none known

Roderick P Venekamp is an Editor for Cochrane Acute Respiratory Infections and Cochrane ENT, but had no role in the editorial process of this review.

Vittoria Lutje: none known.

Eelko Hak has authored a paper on the design of the CAPITA study (*Netherlands Journal of Medicine*), but was not involved in the actual conduct of that study, nor does the study pose a conflict of interest to the current work.

Anne GM Schilder: the evidENT team at University College London is supported in part by the National Institute for Health Research (NIHR) University College London Hospitals Biomedical Research Centre. Our research is funded by the NIHR and EU Horizon2020. I am the national chair of the NIHR Clinical Research Network ENT Specialty. I am the Surgical Specialty Lead for ENT for the Royal College of Surgeons of England's Clinical Trials Initiative. I am co-investigator on the NIHR PGfAR grant 'Defining best Management for Adults with

Chronic Rhinosinusitis: the MACRO Programme'. In my role as director of the NIHR UCLH BRC Deafness and Hearing Problems Theme, I act as an adviser on clinical trial design and delivery to a range of biotech companies.

Elisabeth AM Sanders: for research on pneumococcal vaccines, carriage and surveillance studies, Elisabeth AM Sanders received money paid by governmental agencies and pharmaceutical companies GSK and Pfizer, and paid to the institution or collaborating institutions. Furthermore, Elisabeth AM Sanders has participated in Independent Data Monitoring Committees and Advisory Boards for pharmaceutical companies for vaccine studies and/or respiratory tract infections with fees paid to the institution before 2014. In general, fees were always paid to the institution and used for research purposes.

Roger AMJ Damoiseaux: none known.

Elisabeth AM Sanders and Eelko Hak are authors of studies included in this review (Elisabeth AM Sanders: [Jansen 2008](#); [van Kempen 2006](#); [Veenhoven 2003](#); Eelko Hak: [Jansen 2008](#)). To avoid any potential conflicts of interest, other review authors reviewed the eligibility and performed 'Risk of bias' assessment and data extraction for these studies.

## SOURCES OF SUPPORT

### Internal sources

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### External sources

- No sources of support supplied

## DIFFERENCES BETWEEN PROTOCOL AND REVIEW

Following feedback from a peer reviewer and editorial advice, we changed the title of the 2019 update ([Fortanier 2019](#)) to include the population (children). We also added adverse effects as the co-primary outcome. We created a 'Summary of findings' table for pneumococcal conjugate vaccines (PCVs) administered in early infancy using the following outcomes: frequency of all-cause AOM episodes (co-primary outcome), adverse effects (co-primary outcome), frequency of pneumococcal AOM, and frequency of recurrent AOM (defined as three or more AOM episodes in six months or four or more in one year). Furthermore, we used the five GRADE considerations (study limitations, consistency of effect, imprecision, indirectness, and publication bias) to assess the quality of a body of evidence as it relates to the studies that contribute data to the prespecified outcomes. This was not proposed in the protocol. Also, some (other) Cochrane methods have evolved over time. The revised methods have been applied to this 2020 update where required.

## NOTES

The focus in research has shifted from the use of pneumococcal polysaccharide vaccines (PPVs) to pneumococcal conjugate vaccines (PCVs) in children, and the role of PPVs in the prevention of AOM in children is no longer relevant, as PPVs are no longer used as primary intervention in children since the introduction of PCVs. The focus of the current review has therefore shifted from the effect of PPVs to the effect of PCVs on acute otitis media. No further attention will be paid to the effects of PPVs, which were described in prior versions of this review ([Straetemans 2003](#)).