SYSTEMATIC REVIEW

Global parental acceptance, attitudes, and knowledge regarding human papillomavirus vaccinations for their children: a systematic literature review and metaanalysis

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Abstract

Background This systematic literature review aims to summarize global research on parental acceptance, attitudes, and knowledge regarding human papillomavirus vaccinations.

Methods The literature search was conducted in PubMed, Web of Science and Scopus, and included publications from 2006 to 2023. Study quality was assessed using the Newcastle-Ottawa Scale. The Grading of Recommendations Assessment, Development, and Evaluation guidelines were used to assess the strength of evidence for the primary outcome. Meta-analyses were performed using random-effects models to estimate pooled parental acceptance of HPV vaccinations. Studies were stratified by publication years, and a subgroup analysis was conducted to estimate vaccine acceptance rates by world regions. Additionally, sensitivity analyses examined the role of parents in accepting HPV vaccinations for children of different sexes.

Results Based on 86 studies, we found that parents generally supported HPV vaccinations for their children, yet HPV vaccine acceptance rates showed high variation (12.0 to 97.5%). The subgroup analysis revealed geographical variations in pooled parental HPV vaccine acceptance rates, with the highest rate observed in Africa (79.6%; 95% CI: 73.5–85.2; $I^2 = 98.3\%$; p < 0.01) and the lowest in North America (56.7%; 95% CI: 49.3–64.0; $I^2 = 99.4\%$; p < 0.01). Sensitivity analyses showed that acceptance was higher for daughters than for sons, with mothers more willing to get their daughters vaccinated. The proportion of parents reporting barriers or benefits regarding HPV vaccinations varied widely (0.3 to 95.8%) between study regions. Across all world regions, fear of adverse effects and concerns about vaccine safety were the main barriers, whereas the desire to protect their children from cancer was a significant predictor of vaccine acceptance. Knowledge levels varied widely (6.5 to 100%) between world regions and according to the questions asked. In most studies, knowledge e.g., that HPV is sexually transmitted, and that HPV vaccination provides protection against cervical cancer, ranged from moderate to high.

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Conclusions The results indicated moderate parental acceptance of HPV vaccines. Public knowledge of HPV infection should be promoted, and special efforts should be made to minimize the existing barriers and increase vaccination accessibility and uptake.

Keywords Human papillomavirus, Vaccination, Acceptance, Systematic review, Meta-analysis **PROSPERO registration number** CRD42019135056

Background

HPV and its impact

Human papillomavirus (HPV) is one of the most common sexually transmitted viruses worldwide [1-4]. In their lifetime, most sexually active individuals will be infected at least once, mostly without developing any pathological changes associated with HPV persistence [5–7]. Over 200 types of HPV are known. Low-risk HPV types 6 and 11 are associated with 90% of genital wart cases, while high-risk HPV types 16 and 18 contribute to 70% of all cervical cancer cases [1, 7]. Worldwide, cervical cancer is the fourth most frequent cancer among women, with an estimated 660,000 new cases and 350,000 deaths in 2022 [7]. Almost all cancer cases are caused by HPV. Additionally, HPV infection is associated with the development of the cancers of the head, neck, anus, and genital tract (i.e., penile, vaginal, and vulvar cancers) [8]. Cervical cancer is considered almost completely preventable due to highly effective primary (HPV vaccine) and secondary (screening) prevention measures [9, 10].

Global HPV vaccination efforts

The first HPV vaccine was approved in 2006, marking the beginning of HPV vaccination efforts. Currently, there are six licensed HPV vaccines available: three bivalent (Cervarix[®], Cecolin[®], Walrinvax[®], two quadrivalent (Gardasil°, Cervavax°) and one nonavalent vaccine (Gardasil 9°). Since 2009, four of these vaccines have been prequalified by the World Health Organization (Cecolin°, Cervarix°, Gardasil°, Gardasil 9°). The bivalent vaccine Walrinvax[®] is currently under review by the WHO, while the quadrivalent vaccine Cervavac[®] is licensed for use in specific countries but has not yet received WHO prequalification [11]. The nonavalent HPV vaccine protects against more than 99% of HPV cases related to genotypes 6, 11, 16 and 18 and against up to 96.7% of HPV cases related to genotypes 31, 33, 45, 52, and 58 [12]. As of 2022, 125 countries include HPV vaccine in their routine vaccinations for girls, and 47 countries also for boys [13].

The WHO recommends a one- or two-dose schedule for girls and women between the ages of 9 and 20 years and two doses within a 6-months-interval for women older than 20 years [14]. The strategy of the WHO is to enhance global vaccination programs to increase vaccination rates and incorporate HPV vaccination into national vaccination schemes. As part of this "HPV elimination program", nationwide vaccination rates of 90% are envisioned by 2030 [15].

Global parental acceptance, attitudes, and knowledge

Despite the demonstrated high effectiveness against persistent HPV16 and 18 infections, the parental decision to vaccinate their children remains a matter of debate, and HPV vaccination rates in many countries remain low [16-18]. However, global coverage for the first dose of the HPV vaccine in girls grew from 20% in 2022 to 27% in 2023, indicating some progress in vaccination efforts [19]. Research on HPV vaccine acceptance has primarily focused on mothers and daughters, while little is known about the acceptance of HPV vaccines among parents and their sons [20-22]. Parental attitudes and knowledge significantly influence the acceptance of HPV vaccines worldwide. Positive attitudes towards vaccinations in general and specific trust in the efficacy and safety of HPV vaccines are strongly linked to higher acceptance rates [23–25]. Furthermore, increased knowledge about HPV and its link to cervical cancer enhances vaccine acceptance among parents [26–28].

Previous systematic reviews have aimed at addressing the existing research gaps on knowledge, attitudes, and acceptance rates related to the HPV vaccine. Derbie et al. (2023) and Zewdie et al. (2023) examined Ethiopian parents' attitudes toward vaccinating their children, focusing on local cultural and social factors [29, 30]. Kutz et al. (2023) focused on the awareness and attitudes towards the HPV vaccine in Sub-Saharan Africa, revealing the challenges in promoting vaccination in the region due to limited resources and awareness [31]. López et al. (2020) examined European parental acceptance of HPV vaccines, highlighting the variability in knowledge and acceptance rates across European countries [32]. Suárez et al. (2019) explored the attitudes of Latino fathers in the USA towards the HPV vaccine, pointing out the need for culturally sensitive educational interventions [33]. Perlman et al. (2014) provided a detailed analysis of the knowledge and acceptability of HPV vaccination in Sub-Saharan Africa [34]. Trim et al. (2012) examined parental knowledge and attitudes towards HPV vaccines, emphasizing concerns about safety and information gaps, and how attitudes shifted pre- and post-FDA approval of bivalent and quadrivalent vaccines [35].

Despite these advances in closing knowledge gaps related to HPV vaccines, previous reviews on HPV vaccination had a limited geographic or demographic scope, primarily providing insights into specific regions or populations. This resulted in a lack of a comprehensive global perspective and an incomplete understanding of HPV vaccine acceptance rates, parental attitudes, and knowledge worldwide. For this reason, a comprehensive review and meta-analysis was conducted to qualitatively and quantitatively synthesize existing evidence on parental acceptance, attitudes, and knowledge related to HPV vaccinations for their children. We aimed to (1) quantify the global parental acceptance rates regarding HPV vaccination, (2) identify parental attitudes regarding the perceived benefits and barriers associated with HPV vaccination, and (3) quantify the level of parental knowledge regarding HPV and HPV vaccination.

Methods

This systematic literature review followed a study protocol registered in the International Prospective Register of Systematic Reviews on July 10, 2019 (PROSPERO, CRD42019135056) and adhered to the Preferred Reporting Items in Systematic Reviews and Meta-Analyses statement [36].

Literature search

A systematic literature search for quantitative studies was performed in the literature databases PubMed, Web of Science, and Scopus (Additional file 1). Additionally, a hand search of reference lists from relevant studies was conducted to identify any studies that may have been missed in the database search. Covidence was used to remove duplicates and perform title, abstract, and fulltext screening. Titles and abstracts of the identified studies were screened for inclusion by the lead author (SH). Studies that met the inclusion criteria were forwarded to full-text screening. Two authors (SH, CC) independently reviewed the full texts of the studies. Any discrepancies or disagreements were discussed until a consensus was reached.

Inclusion and exclusion criteria

Since HPV vaccines became available in 2006, quantitative studies published online or on paper in English between January 1, 2006 and October 31, 2023 were included. The inclusion criteria were as follows: (1) parental acceptance rates regarding HPV vaccinations (mandatory); (2) parental attitudes regarding HPV vaccinations (optional); (3) parental knowledge of HPV and HPV vaccinations (optional). In the context of this review, "parental" refers to all persons who have legal and/or factual custody of a child. This includes biological parents, adoptive parents, and foster parents. The study participants included parents and their children eligible for HPV vaccinations.

A study was regarded as measuring HPV vaccination acceptance if it evaluated a positive or negative intention or willingness toward vaccinating children in the future (intention to vaccinate) or having consented (already vaccinated) or not to vaccinate their children in the past.

The systematic review excluded studies based on specific criteria, such as studies examining the acceptance of vaccines against sexually transmitted diseases in general. Reviews and meta-analyses were also excluded. Furthermore, studies conducted or published outside the inclusion period, or with populations not matching the demographic criteria, were not considered. This refers specifically to studies asking adults without children to consider hypothetical children, or those exclusively focusing on a pediatric population, or only involving adults. Studies with inappropriate designs, such as qualitative or interventional studies, and those based on convenience or non-probability sampling, were also not included in this review. Additionally, any study that did not report a parental acceptance rate in percent (%) for HPV vaccination for their children was excluded. Nonresearch materials and studies published in languages other than those specified were also excluded.

Data extraction

Data were extracted by two independent authors (SH, CC). In case of disagreements, a final decision was reached by consensus. The data extracted included study region, year(s) of study conduct, study setting (refers to the specific location or context in which the study was conducted), parents' age and sex, children's age and sex, parents' ethnicity, the total sample size of parents (n), the survey instrument used, acceptance rate (%), type of acceptance, and sampling method.

Quality assessment

Study quality was independently assessed by two authors (SH, VO) using the Newcastle-Ottawa Scale (NOS) for cohort studies and a modified version of the NOS for cross-sectional studies (Additional file 2 and 3). In adapting the NOS for cross-sectional studies, we expanded the rating system from 9 to 10 stars to account for the specific methodological differences of this study design.

Disagreements were discussed until a consensus was reached. Each study was rated "very good," "good," "satisfactory," or "unsatisfactory" based on a star system in which a study is judged according to the study group selection, group comparability, and ascertainment of either the exposure or outcome of interest for studies [37, 38].

The quality of the evidence was evaluated using the Grading of Recommendations Assessment,

Development, and Evaluation (GRADE) criteria (Additional file 4) [39].

Data analysis

We descriptively evaluated parental acceptance, the perceived barriers and benefits of HPV vaccinations, and parental knowledge regarding HPV and HPV vaccinations, and presented the findings in tabular form.

Parental acceptance of HPV vaccinations was categorized according to the type of acceptance (intention to vaccinate, already vaccinated, mixed). The proportions of agreement to seven perceived barriers and benefits (desire to protect their children against cancer, recommendation by a pediatrician or family physician, concern regarding vaccine efficacy, concern regarding the adverse effects of HPV vaccines, fear that vaccination will encourage sexual activity, lack of recommendations, and lack of knowledge) were determined. The knowledge level was determined based on the responses to five key HPV-related questions (aware of HPV, aware of HPV vaccines, aware that HPV is sexually transmitted, aware that cervical cancer is related to HPV infection, and aware that HPV vaccines prevent cervical cancer). We summarized the results by world region: (1) Africa; (2) Asia; (3) Australia; (4) Europe; (5) North America; (6) Oceania; (7) South America, as well as by the sex of parents and the sex of children: (1) Parents and daughters; (2) Parents and sons; (3) Parents and children; (4) Mothers and daughters.

Statistical analysis

A meta-analysis was performed to estimate the overall pooled parental acceptance of HPV vaccination using random-effects models. An additional analysis was conducted to stratify the results by publication year. Only cross-sectional studies that surveyed parents of non-vaccinated children were included. A subgroup analysis was performed based on the world regions. In the sensitivity analyses, we further stratified by the sex of parents and children, as well as by combinations of world region and participant sex.

Statistical heterogeneity among the studies was tested using Cochrane's Q test (significance level p < 0.10). The I² statistic was employed to quantify the heterogeneity of the results using Higgins and Thompson's guidelines, which indicate that I² values of 25% represent low, 50% medium, and 75% represent high heterogeneity [40, 41]. To ensure robustness, additional analyses for publication bias, small-study effects, and overall effect asymmetry were conducted. Publication bias was assessed using a funnel plot (Additional file 5). Additionally, the trim-and-fill method was used, which adjusts for bias by estimating and imputing the number of missing studies needed to achieve symmetry [42, 43]. Egger's regression test was used to evaluate the influence of small studies on the overall effect size, with a significant intercept (p-value < 0.05) indicating potential small-study bias [42].

All data analyses were performed using R^{*} (version 4.0.3), using the metafor package (version 3.0.2).

Results

We identified 4,635 studies in the literature databases: 2,063 from PubMed, 1,884 from Web of Science, and 688 from Scopus (Fig. 1). Through a hand search, we identified 34 additional studies.

After duplicates were removed, 3,078 studies were reviewed for eligibility, based on the titles and abstracts. According to the exclusion criteria, 2,606 studies were excluded. After reading the full texts of the remaining 472 studies, 386 studies were excluded because they did not meet the inclusion criteria. Thus, 86 studies were included in the systematic literature review, and 62 studies were included in the meta-analysis.

Study characteristics

We included 83 cross-sectional and three cohort studies representing data from 251,880 parents (Table 1; Additional file 6). The reported study population consisted primarily of parents and their daughters. Simple random sampling was used in more than half of the studies, followed by clustered sampling and stratified random sampling. Most studies reported on the parents' intention to vaccinate their children. Some studies analyzed the acceptance of HPV vaccinations by parents whose children had been vaccinated at least once. Paper-based questionnaires were used most frequently for data collection, followed by telephone interviews and web-based questionnaires. White parents were the most frequently examined ethnic group, followed by Asians, Blacks, Hispanics and Oceania/Indigenous. The majority of studies included several ethnic groups in their analyses. The studies were mainly conducted in North and South American countries (Canada, USA, Argentina, Brazil). The remaining studies were conducted in European (Austria, Denmark, England, Finland, France, Iceland, Italy, Netherlands, Poland, Romania, Spain, Sweden) and African countries (Ethiopia, Kenya, Morocco, Nigeria, Uganda, Zambia) and the Asia-Pacific Region (China, India, Israel, Korea, Malaysia, Republic of Fiji, Saudi Arabia, Thailand, Vietnam, Australia, New Zealand). Six studies received a study quality rating "very good", 32 studies "good", 44 studies "satisfactory", and four studies "unsatisfactory" (Additional file 7).

Parental acceptance of HPV vaccinations

The HPV vaccine acceptance rates varied widely across the 86 studies (12.0 to 97.5%). Among these, 19 studies reported high acceptance rates (\geq 80%), 62 studies

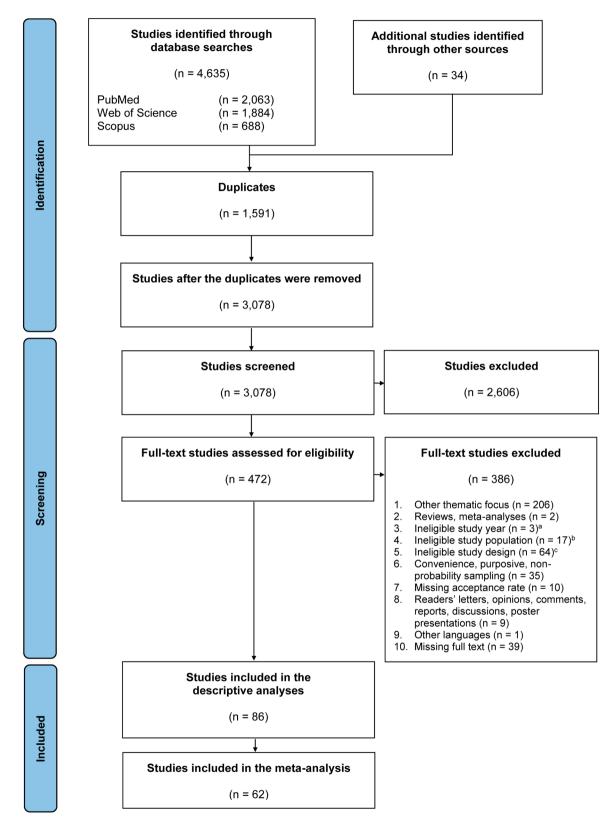


Fig. 1 Flow diagram of the study selection process. ^a Studies that were conducted or published outside the specified inclusion period from January 1, 2006, to August 31, 2023; ^b Studies that did not focus on parents or guardians of children eligible for HPV vaccinations; ^c Studies with inappropriate designs (qualitative studies, interventional studies)

Table 1 Characteristics of the 86 included studies

	Number of studies (n)	Percentage (%)
	86	100
Study design		
Cross-sectional	83	96.51
Cohort	3	3.49
Study participants		
Parents and children	27	31.40
Parents and daughters	31	36.05
Parents and sons	12	13.95
Mothers and daughters	16	18.60
Sample size of the parents (n)		
≤100	2	2.33
101–500	30	34.88
501–999	35	40.70
≥1000	19	22.09
Sampling method		
Stratified random sampling	15	17.44
Systematic sampling	4	4.65
Clustered sampling	17	19.77
Simple random sampling	50	58.14
Year the study was conducted		
2005–2008	23	26.74
2009–2012	23	26.74
2013–2016	17	19.77
2017–2020	9	10.47
2021–2023	8	9.30
Missing	6	6.98
Date of study		
Pre-vaccine licensure ^a	5	5.81
Post-vaccine licensure ^a	81	94.19
Type of acceptance		
Intention to vaccinate	63	73.26
Already vaccinated ^b	12	13.95
Mixed ^c	11	12.79
Survey instrument		
Computer assisted telephone interview (CATI)	3	3.49
Interviewer-administered questionnaire ^d	11	12.79
Interviewer-administered questionnaire (face-to-face interview)	4	4.65
Interviewer-administered questionnaire (telephone interview)	16	18.60
Paper-based questionnaire	22	25.58
Paper-based questionnaire and face-to-face interview	1	1.16
Paper-based questionnaire and telephone interview	3	3.49
Paper-based questionnaire or web-based questionnaire	1	1.16
Self-administered questionnaire ^d	10	11.63
Web-based questionnaire	15	17.44
Study region		
North America	33	38.37
USA	28	32.56
Canada	5	5.81
Africa	19	22,09
Ethiopia	7	8.14
Nigeria	4	4.65
Kenya	2	2.33
Uganda	2	2.33

Table 1 (continued)

	Number of studies (n)	Percentage (%)		
	86	100		
Morocco	1	1.16		
Zambia	1	1.16		
Asia	15	17.44		
China	4	4.65		
India	2	2.33		
Malaysia	2	2.33		
Thailand	2	2.33		
Israel	1	1.16		
Republic of Fiji	1	1.16		
Saudi Arabia	1	1.16		
Korea	1	1.16		
Vietnam	1	1.16		
Europe	15	17.44		
Italy	2	2.33		
Netherlands	2	2.33		
Poland	2	2.33		
Austria	1	1.16		
Denmark	1	1.16		
England	1	1.16		
Finland	1	1.16		
France	1	1.16		
Iceland	1	1.16		
Romania	1	1.16		
Spain	1	1.16		
Sweden	1	1.16		
South America	2	2.33		
Argentina	1	1.16		
Brazil	1	1.16		
Australia	1	1.16		
Oceania	1	1.16		
New Zealand	1	1.16		
Ethnic origin of parents ^e				
White	27	31.40		
Asian	5	5.81		
Black	5	5.81		
Hispanic	5	5.81		
Oceania/Indigenous	1	1.16		
Others	1	1.16		
Missing	42	48.84		
udy quality (based on the NOS)				
Very good	6	6.98		
Good	32	37.21		
Satisfactory	44	51.16		
nsatisfactory	4	4.65		

^a Pre-vaccine licensure: 2006 (study period: 2005), post-vaccine licensure: 2007–2023

 $^{\rm b}$ Children had already been administered at least one dose of the HPV vaccine

 $^{\rm c}$ Intention to vaccinate and already vaccinated

^d No information on format

^e Studies were classified based on the most frequently studied ethnicity

showed moderate acceptance rates (>30% to <80%), and five studies reported low acceptance rates (\leq 30%).

Most of the studies involved parents who intended to vaccinate their children (n=62), while in 12 studies, the vaccine had already been administered. Across all 86 studies, sample sizes varied widely, ranging from 39 to 52,855 parents. Studies with high acceptance rates (\geq 80%) and low acceptance rates (\leq 30%) had smaller sample sizes, ranging from 39 to 1,302 and 368 to 1,255 participants, respectively. No obvious trend toward large or small sample sizes was observed. Additionally, studies reporting low acceptance rates exclusively included parents of both sexes. The age of the parents was higher than that in studies with high acceptance rates. In the 86 studies, acceptance rates were higher for daughters than for sons, with mothers more willing to get their daughters vaccinated.

Meta-analysis

For the meta-analysis, 62 cross-sectional studies that surveyed parents of non-vaccinated children were included (Fig. 2). Acceptance rates varied widely across the 62 studies (28.3 to 94.3%) over the years. The pooled acceptance rate for HPV vaccinations across these studies was 67.2% (95% CI: 62.6–71.7; $I^2 = 99.5\%$; p < 0.01; PI: 30.1–94.8), indicating substantial heterogeneity across studies.

Additional analysis stratified by publication year revealed no association in HPV vaccination acceptance rates (Additional file 8). The visual inspection of the funnel plot showed no asymmetry (Additional file 5). The Trim-and-Fill method indicated that no studies were imputed, suggesting no evidence of asymmetry in the data. Egger's Regression Test for Funnel Plot Asymmetry showed no significant results, confirming the absence of small study bias in the meta-analysis.

Based on the GRADE criteria, the overall vaccine acceptance rate was supported by moderate-quality evidence from 62 studies.

Subgroup analysis by world region

Subgroup analysis by world region revealed that studies from Africa had the highest pooled parental acceptance rate for HPV vaccinations (79.6%; 95% CI: 73.5–85.2; I² = 98.3%; p<0.01; PI: 51.1–97.4), followed by studies from Europe (65.9%; 95% CI: 51.7–78.8; I² = 99.5%; p<0.01; PI: 18.8–98.5), Asia (63.7%; 95% CI: 55.4–71.6; I² = 99.1%; p<0.01; PI: 32.8–89.4), and North America (56.7%; 95% CI: 49.3–64.0; I² = 99.4%; p<0.01; PI: 18.8–98.5) (Fig. 3).

The quality of evidence varied across subgroups. For African studies, moderate evidence was observed, while lower quality evidence was found for Asian and North American studies. Studies from Europe presented very low-quality evidence, with a wide PI, highlighting significant uncertainty. Page 8 of 22

Sensitivity analyses by sex of parents and their children

The sensitivity analysis by the sex of the parents and their children revealed that the highest pooled acceptance rate was among mothers of daughters (73.9%; 95% CI: 65.7–81.3%; $I^2 = 99.1\%$; p < 0.01; PI: 41.8–95.9), and the lowest was among parents of sons (57.7%; 95% CI: 47.7–67.5%; $I^2 = 98.6\%$; p < 0.01; PI: 25.7–86.5) (Fig. 4). In an analysis stratified by both sex and world regions, the highest rate was among mothers and daughters in studies from Africa (86.3%; 95% CI: 81.6–90.5; $I^2 = 89.4\%$; p < 0.01; PI: 75.5–94.9), and the lowest was among parents and sons in North American studies (51.2%; 95% CI: 42.2–60.2; $I^2 = 97.1\%$; p < 0.01; PI: 29.9–72.3) (Additional file 9).

Sensitivity analyses further underscored challenges, particularly in the "Parents and sons" subgroup, which had very low-quality evidence due to small studies with low study quality. In some cases, publication bias reduced the reliability of findings, notably in the "Parents and daughters, North America", "Parents and sons, Europe" and "Mothers and daughters, North America" subgroups, which were also impacted by small studies and low study quality.

Parental attitudes regarding HPV and HPV vaccinations

We extracted data for seven parental barriers and benefits from 43 studies conducted in Africa (n=12), Asia (n=6), Australia (n=1), Europe (n=11), North America (n=10), Oceania (n=1) and South America (n=2) (Table 2).

In total, 181,736 parents aged 18 to 82 years reported one or several perceived barriers and benefits for parental vaccination intention. Four studies had a study quality rating of "very good", 14 studies were "good", 23 studies were "satisfactory", and two studies were "unsatisfactory".

The most frequently cited benefit in the studies was the desire to protect children from cancer, while the most frequently cited barrier was concern about the adverse effects of the HPV vaccine.

Regarding benefits, the highest proportion was observed in a study among parents of daughters in North America (USA) and the lowest was noted among European parents of children (Sweden). Regarding barriers, the highest proportion was observed in a study among European parents of children (Sweden) and the lowest was observed among North American parents of children (USA).

Benefit 1: Desire to protect their children against cancer

In African studies (Ethiopia; n=2), 40.4 to 77.0% of parents reported cancer protection as a benefit of HPV vaccinations [44, 45], which is higher compared to European studies (Denmark, Italy, Sweden; n=3), where the perception ranged from 6.0 to 67.0% [46–48]. In Asia (India, Saudi Arabia; n=3), the perception of cancer protection as a benefit of HPV vaccination was reported by

Study	Total	Events		Acceptance Rate [95% CI]
Wang L.D-L. et al., 2015	368	104	├─■ ─┤	0.283 [0.238, 0.330
Calo et al., 2017	1255	360	┝╼┤	0.287 [0.262, 0.312
Borena et al., 2016	148	48		0.324 [0.252, 0.402
Voidăzan et al., 2016	918	327	├─ ● ─┤ .	0.356 [0.326, 0.387
Litton et al., 2011	403	150		0.372 [0.326, 0.420
Huon et al., 2020	127	48		0.378 [0.296, 0.464]
Humnesa et al., 2022	619	249		0.402 [0.364, 0.441]
Cheruvu et al., 2017	21467	8751		0.408 [0.401, 0.414]
Moss et al., 2015 Clark et al., 2016	412	177		0.430 [0.382, 0.478] 0.433 [0.398, 0.469]
Songthap et al., 2012	734 648	318 290		0.448 [0.409, 0.486]
van Keulen et al., 2013	952	428		0.450 [0.418, 0.481]
Askelson et al., 2010	217	104		0.479 [0.413, 0.546]
Nguyen et al., 2022	785	386		0.492 [0.457, 0.527]
McRee et al., 2013	506	254		0.502 [0.458, 0.545]
Dempsey et al., 2011	1178	601		0.510 [0.482, 0.539]
Mansfield et al., 2018	1037	585		0.564 [0.534, 0.594]
Wang Z. et al., 2018	296	167		0.564 [0.507, 0.620]
Pourat et al., 2012	4896	2766	·	0.565 [0.551, 0.579]
Fang et al., 2010	1383	795	⊢∎⊣	0.575 [0.549, 0.601]
M'Imunya et al., 2011	332	193		0.581 [0.528, 0.634]
Guerry et al., 2011	387	240		0.620 [0.571, 0.668]
Mohd Sopian et al., 2018	280	175	⊢_ ∎	0.625 [0.568, 0.681]
Dahlström et al., 2010	13840	8691		0.628 [0.620, 0.636]
Shibli et al., 2019	313	206		0.658 [0.605, 0.710]
Lin et al., 2019	5799	3842		0.663 [0.650, 0.675]
Kadis et al., 2011	496	332	⊢−■−−┤	0.669 [0.627, 0.710]
Rose et al., 2012	769	515		0.670 [0.636, 0.702]
Hopenhayn et al., 2007	626	423		0.676 [0.639, 0.712]
Ogilvie et al., 2008	1381	936		0.678 [0.653, 0.702]
Bianco et al., 2014 Madhivanan et al., 2014	566	402		0.710 [0.672, 0.747] 0.715 [0.682, 0.746]
Rabiu et al., 2020	778	556		0.720 [0.670, 0.768]
Selmouni et al., 2015	318 1312	229 956		0.729 [0.704, 0.752]
Lin et al., 2020	5702	4213		0.739 [0.727, 0.750]
Arrossi et al., 2012	100	74		0.740 [0.650, 0.821]
Kruiroongroj et al., 2014	758	564	'⊢∎→	0.744 [0.712, 0.774
Ezat et al., 2013	155	116		0.748 [0.677, 0.813]
Constantine et al., 2007	522	392	`⊢-∎ `	0.751 [0.713, 0.787
Oh et al., 2010	1000	751	· · · · · · · · · · · · · · · · · · ·	0.751 [0.724, 0.777]
Lai et al., 2013	804	606	⊢_∎	0.754 [0.723, 0.783]
Dairo et al., 2016	612	466	┝╼┻─┤	0.761 [0.727, 0.794]
Marshall et al., 2007	601	462	⊢ ∎	0.769 [0.734, 0.802]
Sinshaw et al., 2022	601	465		0.774 [0.739, 0.806]
Muhwezi et al., 2014	870	681		0.783 [0.755, 0.810]
Aragaw et al., 2023	721	570		0.791 [0.760, 0.819]
Destaw et al., 2021	502	399		0.795 [0.758, 0.829]
Degarege et al., 2018	831	664		0.799 [0.771, 0.826]
Mortensen et al., 2010 Brabin et al., 2006	450	360		0.800 [0.762, 0.836] 0.811 [0.766, 0.852]
Alene et al., 2020	317	257		0.813 [0.787, 0.838]
Larebo et al., 2020	899 530	731 450		0.849 [0.817, 0.878]
Ganczak et al., 2018	450	450 383		0.851 [0.817, 0.882]
Ezenwa et al., 2013	450 290	248		0.855 [0.812, 0.893]
Woodhall et al., 2007	727	622		0.856 [0.829, 0.880]
Bernat et al., 2009	1504	1302	,	0.866 [0.848, 0.882]
Vermandere et al., 2014	287	253		0.882 [0.842, 0.916]
Morhason-Bello et al., 2015	1002	888	, - , ⊢ ∎⊣	0.886 [0.866, 0.905]
Azuogu et al., 2019	267	238		0.891 [0.851, 0.926]
Ndejjo et al., 2017	900	813	· ⊢∎-í	0.903 [0.883, 0.922]
Oddsson et al., 2016	583	530	⊢∎-1	0.909 [0.884, 0.931]
Dereje et al., 2021	422	398	. ⊢•+	0.943 [0.919, 0.963]
RE Model of subgroup (Q = 103 Prediction Model	308.24,d	f = 61 , p <	.0.01; l ² = 99.5 %)	0.672 [0.626, 0.717 [0.301, 0.948]
calouon model				
			· · ·	
	0.000		0.250 0.500 0.750	1.000

Fig. 2 Forest plot for HPV vaccine acceptance rates among parents and their children showing individual study estimates and pooled estimate (random-effects model) (n=62). CI=Confidence interval; df=Degrees of freedom; Events=The number of parents/guardians in each study who reported acceptance of the HPV vaccine for their children; l^2 = I-squared statistic, indicating the percentage of variation due to heterogeneity; p < 0.01 = p-value, indicating statistical significance; Q=Cochran's Q statistic for heterogeneity; RE Model=Random-effects model; Total=The total number of parents/guardians in each study included in the meta-analysis

Africa	Total Events		Acceptance Rate [95% C
lumnesa et al., 2022	619 249		0.402 [0.364, 0.441
//Imunya et al., 2011 Rabiu et al., 2020	332 193 318 229		0.581 [0.528, 0.634 0.720 [0.670, 0.768
elmouni et al., 2015 Dairo et al., 2016	1312 956 612 466		0.729 [0.704, 0.752 0.761 [0.727, 0.794
Sinshaw et al., 2022	601 465		0.774 [0.739, 0.806
/uhwezi et al., 2014	870 681		0.783 [0.755, 0.810
Aragaw et al., 2023	721 570		0.791 [0.760, 0.819
Destaw et al., 2021	502 399		0.795 [0.758, 0.829
lene et al., 2020	899 731		0.813 [0.787, 0.838
arebo et al., 2022	530 450	·	0.849 [0.817, 0.878
Ezenwa et al., 2013	290 248		0.855 [0.812, 0.893
Vermandere et al., 2014	287 253		0.882 0.842, 0.916
Morhason-Bello et al., 2015	1002 888	·. ⊢ = ⊣`.	0.886 [0.866, 0.905
Azuogu et al., 2019	267 238		0.891 [0.851, 0.926
Ndejjo et al., 2017	900 813		0.903 [0.883, 0.922
Dereje et al., 2021	422 398		0.943 [0.919, 0.963
Prediction Model	19, df = 16, p < 0.01; l ² = 98		0.796 [0.735, 0.852 [0.511 , 0.974]
Asia Vang L.D-L. et al., 2015	368 104		0.283 [0.238, 0.330
Songthap et al., 2012	648 290		0.448 [0.409, 0.486
Nguyen et al., 2022	785 386		0.492 0.457, 0.527
Wang Z. et al., 2018	296 167		0.564 [0.507, 0.620
Mohd Sopian et al., 2018	280 175	· ⊢_ `	0.625 [0.568, 0.681
Shibli et al., 2019	313 206	·	0.658 [0.605, 0.710
Lin et al., 2019	5799 3842	· • • • • • • • • • • • • • • • • • • •	0.663 [0.650, 0.675
Madhivanan et al., 2014	778 556		0.715 [0.682, 0.746
Lin et al., 2020	5702 4213		0.739 [0.727, 0.750
Kruiroongroj et al., 2014	758 564	, ⊢■-1 ,	0.744 [0.712, 0.774
Ezat et al., 2013 Ob et al. 2010	155 116		0.748 [0.677, 0.813 0.751 [0.724, 0.777
Oh et al., 2010 Degarege et al., 2018	1000 751 831 664		0.799 [0.771, 0.826
RE Model of subgroup (Q = 742 Prediction Model	2.87 , df = 12 , p < 0.01 ; l ² = 99	<i>9</i> ,1 %)	0.637 [0.554, 0.716 [0.328 , 0.894]
Australia Marshall et al., 2007	601 462		0.769 [0.734, 0.802
Europe			
Borena et al., 2016	148 48		0.324 [0.252, 0.402
Voidăzan et al., 2016	918 327	,	0.356 0.326, 0.387
Huon et al., 2020 van Keulen et al., 2013	127 48		0.378 [0.296, 0.464
Dahlström et al., 2010	952 428 13840 8691		0.450 [0.418, 0.481 0.628 [0.620, 0.636
Bianco et al., 2014	566 402		0.710 [0.672, 0.747
Mortensen et al., 2010	450 360		0.800 [0.762, 0.836
Brabin et al., 2006	317 257		0.811 [0.766, 0.852
Ganczak et al., 2018	450 383		0.851 [0.817, 0.882
Woodhall et al., 2007	727 622	· · · · · · · · · · · · · · · · · · ·	0.856 0.829, 0.880
Oddsson et al., 2016	583 530	╵╴┝╼╌┤	0.909 [0.884, 0.931
RE Model of subgroup (Q = 123 Prediction Model	1.27, df = 10, p < 0.01; l ² = 9	99.5 %)	0.659 [0.517, 0.788 [0.188 , 0.985]
North America			
Calo et al., 2017	1255 360		0.287 [0.262, 0.312
	403 150		0.372 [0.326, 0.420
Litton et al., 2011			
Litton et al., 2011 Cheruvu et al., 2017	21467 8751		0.408 [0.401, 0.41
Litton et al., 2011 Cheruvu et al., 2017 Moss et al., 2015	21467 8751 412 177		0.408 [0.401, 0.414 0.430 [0.382, 0.478
Litton et al., 2011 Cheruvu et al., 2017 Moss et al., 2015 Clark et al., 2016	21467 8751 412 177 734 318		0.408 [0.401, 0.41 0.430 [0.382, 0.47 0.433 [0.398, 0.46
Litton et al., 2011 Cheruvu et al., 2017 Moss et al., 2015 Clark et al., 2016 Askelson et al., 2010	21467 8751 412 177 734 318 217 104		0.408 0.401, 0.41 0.430 0.382, 0.47 0.433 0.398, 0.46 0.479 0.413, 0.54
Litton et al., 2011 Cheruvu et al., 2017 Moss et al., 2015 Clark et al., 2016 Askelson et al., 2010 McRee et al., 2013 Dempsey et al., 2011	21467 8751 412 177 734 318		0.408 [0.401, 0.41 0.430 [0.382, 0.47 0.433 [0.398, 0.46 0.479 [0.413, 0.54 0.502 [0.458, 0.54 0.510 [0.482, 0.53
Litton et al., 2011 Cheruvu et al., 2017 Moss et al., 2015 Clark et al., 2016 Askelson et al., 2010 WcRee et al., 2013 Dempsey et al., 2011 Mansfield et al., 2018	21467 8751 412 177 734 318 217 104 506 254		0.408 [0.401, 0.41 0.430 [0.382, 0.47 0.433 [0.398, 0.46 0.479 [0.413, 0.54 0.502 [0.458, 0.54 0.510 [0.482, 0.53
Litton et al., 2011 Cheruvu et al., 2017 Moss et al., 2015 Clark et al., 2016 Askelson et al., 2010 McRee et al., 2013 Dempsey et al., 2011 Mansfield et al., 2018 Pourat et al., 2012	21467 8751 412 177 734 318 217 104 506 254 1178 601 1037 585 4896 2766		$\begin{array}{c} 0.408 \left[0.401, 0.41 \right. \\ 0.430 \left[0.382, 0.47 \right. \\ 0.433 \left[0.398, 0.46 \right. \\ 0.479 \left[0.413, 0.54 \right. \\ 0.502 \left[0.458, 0.54 \right. \\ 0.510 \left[0.482, 0.53 \right. \\ 0.564 \left[0.534, 0.59 \right. \\ 0.565 \left[0.551, 0.57 \right. \end{array} \right]$
Litton et al., 2011 Cheruvu et al., 2017 Moss et al., 2015 Clark et al., 2016 Askelson et al., 2010 McRee et al., 2010 Dempsey et al., 2011 Mansfield et al., 2018 Pourat et al., 2010 Fang et al., 2010	21467 8751 412 177 734 318 217 104 506 254 1178 601 1037 585 4896 2766 1383 795		0.408 [0.401, 0.41 0.430 [0.382, 0.47 0.433 [0.398, 0.46 0.479 [0.413, 0.54 0.510 [0.482, 0.53 0.564 [0.534, 0.59 0.565 [0.551, 0.57 0.575 [0.549, 0.60
Litton et al., 2011 Chenvux et al., 2017 Voss et al., 2015 Clark et al., 2016 Askelson et al., 2010 MeRee et al., 2010 Dempsey et al., 2011 Mansfield et al., 2018 Pourat et al., 2012 Fang et al., 2010 Guerry et al., 2011	21467 8751 412 177 734 318 217 104 506 254 1178 601 1037 585 4896 2766 1383 795 387 240		$\begin{array}{c} 0.408 \left[0.401 , 0.41 \right] \\ 0.430 \left[0.382 , 0.47 \right] \\ 0.433 \left[0.382 , 0.47 \right] \\ 0.479 \left[0.413 , 0.54 \right] \\ 0.502 \left[0.458 , 0.54 \right] \\ 0.510 \left[0.482 , 0.53 \right] \\ 0.564 \left[0.534 , 0.59 \right] \\ 0.565 \left[0.551 , 0.57 \right] \\ 0.575 \left[0.549 , 0.60 \right] \\ 0.620 \left[0.571 , 1 \right] \\ 0.64 \left[0.534 \right] \\ 0.56 \left[0.551 , 0.57 \right] \\ 0.575 \left[0.549 , 0.60 \right] \\ 0.620 \left[0.571 , 1 \right] \\ 0.68 \left[0.534 \right] \\ 0.58 \left[0.534 \right] \\ 0.58$
Litton et al., 2011 Cheruvu et al., 2017 Moss et al., 2015 Sakelson et al., 2010 McRee et al., 2010 Dempsey et al., 2011 Pampsey et al., 2011 Guerry et al., 2010 Guerry et al., 2011 Guisert al., 2011	21467 8751 412 177 734 318 217 104 506 254 1178 601 1037 585 4896 2766 1383 795 387 240 496 332		0.408 [0.401, 0.41 0.430 [0.382, 0.47 0.433 [0.398, 0.46 0.479 [0.413, 0.54 0.502 [0.458, 0.54 0.510 [0.482, 0.53 0.564 [0.534, 0.59 0.565 [0.551, 0.57 0.575 [0.549, 0.60 0.629 [0.527, 0.71, 0.66
Litton et al., 2011 Cheruvu et al., 2017 Moss et al., 2015 Clark et al., 2016 Askelson et al., 2010 Dempsey et al., 2010 Dempsey et al., 2011 Mansfield et al., 2018 Pourat et al., 2012 Guerry et al., 2011 Kadis et al., 2011 Hopenhayn et al., 2017	$\begin{array}{rrrr} 21467 & 8751 \\ 412 & 177 \\ 734 & 318 \\ 217 & 104 \\ 506 & 254 \\ 1178 & 601 \\ 1037 & 585 \\ 4896 & 2766 \\ 1383 & 795 \\ 387 & 240 \\ 496 & 332 \\ 626 & 423 \\ \end{array}$		0.408 [0.401, 0.41 0.430 [0.382, 0.47 0.433 [0.398, 0.46 0.570 [0.458, 0.54 0.550 [0.458, 0.54 0.565 [0.551, 0.57 0.565 [0.551, 0.57 0.565 [0.554, 0.60 0.620 [0.571, 0.66 0.669 [0.639, 0.71
Litton et al., 2011 Cheruvu et al., 2017 Moss et al., 2015 Sakkelson et al., 2010 McRee et al., 2010 Dempsey et al., 2011 Pourat et al., 2011 Fang et al., 2011 Guerry et al., 2011 Gadie et al., 2011 Hopenhayn et al., 2007 Oglivie et al., 2008	21467 8751 412 177 734 318 217 104 506 254 1178 601 1037 585 4896 2766 1383 795 387 240 496 332 626 423 1381 936		$\begin{array}{c} 0.408 \left[0.401 \right], 0.41 \\ 0.430 \left[0.382 \right], 0.47 \\ 0.433 \left[0.388 \right], 0.47 \\ 0.433 \left[0.388 \right], 0.47 \\ 0.502 \left[0.458 \right], 0.543 \\ 0.502 \left[0.458 \right], 0.534 \\ 0.550 \left[0.458 \right], 0.545 \\ 0.551 \left[0.575 \right], 0.575 \\ 0.575 \left[0.575 \right], 0.575 \\ 0.575 \left[0.549 \right], 0.66 \\ 0.620 \left[0.577 \right], 1, 0.66 \\ 0.668 \left[0.623 \right], 0.71 \\ 0.676 \left[0.633 \right], 0.71 \\ 0.676 \left[0.633 \right], 0.71 \\ 0.678 \left[0.653 \right], 0.77 \\ 0.575 \left[0.553 \right], 0.71 \\ 0.678 \left[0.553 \right], 0.71 \\ 0.678 \left[0.553 \right], 0.71 \\ 0.578 \left[0.558 \right], 0.578 \\ 0.578 \left[0.558 \right], 0.578 \\ 0.578 \left[0.578 \right], 0.578$
Litton et al., 2011 Cheruvu et al., 2017 Moss et al., 2015 Sakelson et al., 2010 McRee et al., 2010 McRee et al., 2010 Dempsey et al., 2011 Mansfield et al., 2018 Pourat et al., 2011 Guerry et al., 2011 Guerry et al., 2011 Hopenhayn et al., 2007 Dglivie et al., 2007	21467 8751 412 177 734 318 217 104 506 254 1178 601 1037 585 4896 2766 1383 795 387 240 496 332 626 423 1381 936 522 392		$\begin{array}{c} 0.408 \\ [0.401, 0.41, 0.41, 0.430 \\ 0.438 \\ [0.472] \\ 0.473 \\ [0.458, 0.474 \\ 0.552 \\ [0.458, 0.544 \\ 0.552 \\ [0.458, 0.544 \\ 0.552 \\ [0.554, 0.557 \\ 0.575 \\ 0.575 \\ [0.571, 0.586 \\ 0.669 \\ [0.627, 0.711 \\ 0.678 \\ [0.638, 0.757 \\ 0.757 \\ [0.538, 0.707 \\ 0.757 \\ [0.538, 0.707 \\ 0.757 \\ [0.538, 0.707 \\ 0.757 \\ [0.538, 0.707 \\ 0.757 \\ [0.538, 0.707 \\ 0.751 \\ [0.713, 0.781 \\ 0.751 \\ [0.713, 0.781 \\ 0.751 \\ [0.713, 0.781 \\ 0.751 \\ 0.751 \\ [0.713, 0.781 \\ 0.751 \\ 0$
Litton et al., 2011 Cheruvu et al., 2017 Moss et al., 2015 Slark et al., 2016 Skeksion et al., 2010 McRee et al., 2010 Dempsey et al., 2011 Mansfield et al., 2011 Pourat et al., 2012 Eang et al., 2010 Guerry et al., 2011 Kadis et al., 2011 Hopenhayn et al., 2007 Jojikie et al., 2008 Constantine et al., 2007 Lai et al., 2013	21467 8751 412 177 734 318 217 104 506 254 1178 601 1037 585 4896 2766 1383 795 387 240 496 332 626 423 1381 936		$\begin{array}{c} 0.408 \left[0.401, 0.414 \\ 0.430 \left[0.332, 0.475 \\ 0.433 \left[0.339, 0.465 \\ 0.532 \left[0.473 \right] 0.413, 0.546 \\ 0.502 \left[0.458, 0.545 \\ 0.510 \left[0.482, 0.535 \\ 0.565 \left[0.551 \right] 0.577 \\ 0.575 \left[0.549 \right] 0.605 \\ 0.665 \left[0.551 \right] 0.577 \\ 0.678 \left[0.634 \right] 0.610 \\ 0.620 \left[0.627 \right] 0.71 \\ 0.678 \left[0.653 \right] 0.72 \\ 0.75 \left[0.751 \right] 0.566 \\ 0.669 \left[0.627 \right] 0.71 \\ 0.678 \left[0.653 \right] 0.72 \\ 0.75 \left[0.731 \right] 0.78 \\ 0.75 \left[0.731 \right] 0.78 \\ 0.75 \left[0.731 \right] 0.78 \\ 0.75 \left[0.723 \right] 0.732 \\ 0.75 \left[0.723 \right] 0.75 \\ 0.75 \left[0.723 \left] 0.732 \\ 0.75 \left[0.723 \right] 0.732 \\ 0.75 \left[0.723 \left] 0.732 \\ 0.75 \left[0.723 \right] 0.75 \\ 0.75 \left[0.723 \left] 0.725 \\ 0.75 \left[0.723 \left] 0.725 \\ 0.75 \left[0.723 \left] 0.75 \\ 0.75 \left[0.75 \right] 0.75 \\ 0.75 \left[0.75 \left] 0.75 \\ 0.75 \left[0.75 \left] 0.75 \\ 0.75 \left[0.75 \left] 0.75 \\ 0.$
Litton et al., 2011 Cheruvu et al., 2017 Moss et al., 2015 Clark et al., 2016 Askelson et al., 2010 Dempsey et al., 2011 Dempsey et al., 2011 Mansfield et al., 2011 Fang et al., 2010 Guerry et al., 2010 Guerry et al., 2011 Hopenhayn et al., 2007 Oglivie et al., 2008 Constantine et al., 2007 Lai et al., 2009 RE Model of subgroup (Q = 287	$\begin{array}{rrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrr$		$\begin{array}{c} 0.408 \left[0.401 , 0.414 \\ 0.430 \left[0.382 , 0.474 \\ 0.433 \right] 0.398 , 0.476 \\ 0.479 \left[0.413 , 0.546 \\ 0.502 \left[0.458 , 0.544 \\ 0.502 \left[0.458 , 0.544 \\ 0.502 \left[0.548 , 0.594 \\ 0.565 \right] 0.551 \\ 0.575 \left[0.573 \\ 0.575 \\ 0.575 \\ 0.575 \\ 0.575 \\ 0.575 \\ 0.575 \\ 0.751 \\ 0.751 \\ 0.751 \\ 0.751 \\ 0.751 \\ 0.751 \\ 0.751 \\ 0.751 \\ 0.751 \\ 0.756 \\ 0.680 $
Litton et al., 2011 Cheruvu et al., 2017 Moss et al., 2015 Clark et al., 2016 Askelson et al., 2010 McRee et al., 2010 Dempsey et al., 2011 Mansfield et al., 2011 Fang et al., 2012 Fang et al., 2010 Guerry et al., 2011 Hopenhayn et al., 2007 Ogliki et al., 2008 Constantine et al., 2007 Constantine et al., 2007 Lai et al., 2008 Bernat et al., 2009 RE Model of subgroup (Q = 287 Prediction Model	$\begin{array}{rrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrr$		$\begin{array}{c} 0.408 \begin{bmatrix} 0.401 \\ 0.382 \\ 0.382 \\ 0.438 \\ [0.382 \\ 0.478 \\ 0.479 \\ [0.473 \\ 0.479 \\ 0.479 \\ 0.479 \\ 0.479 \\ 0.479 \\ 0.565 \\ 0.551 \\ 0.54 \\ 0.544 \\ 0.551 \\ 0.571 \\ 0.678 \\ 0.678 \\ 0.678 \\ 0.678 \\ 0.678 \\ 0.678 \\ 0.754 \\ 0.723 \\ 0.784 \\ 0.478 \\ 0.567 \\ 0.493 \\ 0.640 \\ 0.258 \\ 0.685 \\ 0.493 \\ 0.640 \\ 0.258 \\ 0.685 \\ 0.258 \\ 0.567 \\ 0.493 \\ 0.640 \\ 0.258 \\ 0.856 \\ 0.885 \\ 0.258 \\ 0.856 \\ 0.885 \\ 0.567 \\ 0.493 \\ 0.640 \\ 0.258 \\ 0.856 \\ 0.885 $
Litton et al., 2011 Cheruvu et al., 2017 Moss et al., 2015 Clark et al., 2016 Askelson et al., 2010 McRee et al., 2010 Dempsey et al., 2011 Pourat et al., 2012 Fang et al., 2012 Guerry et al., 2011 Madis et al., 2011 Mogenhayn et al., 2007 Oglivie et al., 2008 Constantine et al., 2007 at et al., 2013 Bernat et al., 2009 RE Model of subgroup (Q = 287 Prediction Model	$\begin{array}{rrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrr$		$\begin{array}{c} 0.408 \\ [0.401, 0.382, 0.474 \\ 0.433 \\ [0.382, 0.475 \\ 0.475 \\ [0.478, 0.443 \\ 0.502 \\ [0.478, 0.544 \\ 0.502 \\ [0.458, 0.544 \\ 0.554 \\ 0.554 \\ 0.554 \\ 0.554 \\ 0.554 \\ 0.554 \\ 0.554 \\ 0.554 \\ 0.554 \\ 0.554 \\ 0.554 \\ 0.554 \\ 0.554 \\ 0.554 \\ 0.554 \\ 0.554 \\ 0.554 \\ 0.554 \\ 0.555 \\ 0.575 \\ 0.575 \\ 0.575 \\ 0.575 \\ 0.575 \\ 0.575 \\ 0.575 \\ 0.575 \\ 0.575 \\ 0.575 \\ 0.575 \\ 0.575 \\ 0.583 \\ 0.565 \\ 0.483 \\ 0.685 \\ $
Litton et al., 2011 Cheruvu et al., 2017 Moss et al., 2015 Glark et al., 2016 Askelson et al., 2010 McRee et al., 2010 Dempsey et al., 2011 Dempsey et al., 2011 Mansfield et al., 2012 Fang et al., 2012 Guerry et al., 2011 Kadis et al., 2011 Mogenhayn et al., 2007 Ogilivie et al., 2007 Gonstantine et al., 2007 Lai et al., 2013 Bernat et al., 2009 RE Model of subgroup (Q = 287 Prediction Model Oceania Rose et al., 2012	$\begin{array}{ccccc} 21467 & 8751 \\ 412 & 177 \\ 734 & 318 \\ 217 & 104 \\ 506 & 254 \\ 1178 & 601 \\ 1037 & 585 \\ 4896 & 2766 \\ 1383 & 795 \\ 387 & 240 \\ 496 & 332 \\ 626 & 423 \\ 1381 & 936 \\ 522 & 392 \\ 804 & 606 \\ 1504 & 1302 \\ 4.67 , df = 17 , p < 0.01 ; l^2 = 9 \\ 769 & 515 \end{array}$		$\begin{array}{c} 0.406 \left[0.401, 0.414 \\ 0.430 \left[0.382, 0.478 \\ 0.432 \left[0.388 \right] 0.478 \\ 0.433 \left[0.388 \right] 0.478 \\ 0.512 \left[0.458 , 0.545 \\ 0.510 \left[0.482 , 0.539 \\ 0.564 \left[0.554 \right] 0.594 \\ 0.565 \left[0.551 , 0.579 \\ 0.575 \left[0.579 , 0.610 \\ 0.676 \left[0.639 , 0.712 \\ 0.676 \left[0.639 , 0.712 \\ 0.751 \left[0.713 , 0.770 \\ 0.754 \left[0.633 , 0.742 \\ 0.565 \right] 0.549 \\ 0.666 \left[0.648 \right] 0.882 \\ 0.567 \left[0.493 , 0.640 \\ 0.657 \left[0.493 , 0.640 \\ 0.657 \left[0.636 , 0.702 \\ 0.570 \left[0.636 , 0.702 \right] \right] \\ \end{array} \right.$
Litton et al., 2011 Cheruvu et al., 2017 Moss et al., 2015 Clark et al., 2016 Askelson et al., 2010 Dempsey et al., 2011 Dempsey et al., 2011 Dempsey et al., 2011 Mansfield et al., 2011 Fang et al., 2010 Guerry et al., 2010 Guerry et al., 2010 Goglivie et al., 2007 Constantine et al., 2007 Constantine et al., 2007 Lai et al., 2013 Bernat et al., 2009 RE Model of subgroup (Q = 287 Prediction Model Oceania Rose et al., 2012	$\begin{array}{rrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrr$		0.408 [0.401, 0.414 0.430 [0.382, 0.474 0.433 [0.398, 0.466 0.479 [0.413, 0.546 0.502 [0.458, 0.544 0.510 [0.482, 0.533 0.564 [0.534, 0.534 0.556 [0.551, 0.573 0.575 [0.549, 0.601 0.620 [0.571, 0.666 0.669 [0.627, 0.710 0.676 [0.639, 0.712 0.676 [0.633, 0.702 0.751 [0.713, 0.787 0.754 [0.723, 0.783 0.866 [0.6484, 0.882 [0.256 , 0.485] 0.670 [0.636, 0.702
Litton et al., 2011 Cheruvu et al., 2017 Voss et al., 2015 Clark et al., 2016 Askelson et al., 2010 VorBee et al., 2010 Dempsey et al., 2011 Dempsey et al., 2011 Pourat et al., 2012 Fang et al., 2012 Guerry et al., 2011 Kadis et al., 2011 Vogenhayn et al., 2007 Oglivie et al., 2007 Oglivie et al., 2008 Constantine et al., 2007 ARE Model of subgroup (Q = 287 Prediction Model Docenia Rose et al., 2012 South America	$\begin{array}{ccccc} 21467 & 8751 \\ 412 & 177 \\ 734 & 318 \\ 217 & 104 \\ 506 & 254 \\ 1178 & 601 \\ 1037 & 585 \\ 4896 & 2766 \\ 1383 & 795 \\ 387 & 240 \\ 496 & 332 \\ 626 & 423 \\ 1381 & 936 \\ 522 & 392 \\ 804 & 606 \\ 1504 & 1302 \\ 4.67 , df = 17 , p < 0.01 ; l^2 = 9 \\ 769 & 515 \end{array}$		0.408 [0.401, 0.414 0.430 [0.382, 0.474 0.433 [0.398, 0.466 0.479 [0.413, 0.546 0.502 [0.458, 0.544 0.510 [0.482, 0.533 0.564 [0.534, 0.534 0.556 [0.551, 0.573 0.575 [0.549, 0.601 0.620 [0.571, 0.666 0.669 [0.627, 0.710 0.676 [0.639, 0.712 0.676 [0.633, 0.702 0.751 [0.713, 0.787 0.754 [0.723, 0.783 0.866 [0.6484, 0.882 [0.256 , 0.485] 0.670 [0.636, 0.702
Litton et al., 2011 Cheruvu et al., 2017 Moss et al., 2015 Glark et al., 2016 Askelson et al., 2010 McRee et al., 2010 Dempsey et al., 2011 Dempsey et al., 2011 Mansfield et al., 2012 Fang et al., 2012 Guerry et al., 2011 Kadis et al., 2011 Mogenhayn et al., 2007 Ogilivie et al., 2007 Gonstantine et al., 2007 Lai et al., 2013 Bernat et al., 2009 RE Model of subgroup (Q = 287 Prediction Model Oceania Rose et al., 2012	$\begin{array}{ccccc} 21467 & 8751 \\ 412 & 177 \\ 734 & 318 \\ 217 & 104 \\ 506 & 254 \\ 1178 & 601 \\ 1037 & 585 \\ 4896 & 2766 \\ 1383 & 795 \\ 387 & 240 \\ 496 & 332 \\ 626 & 423 \\ 1381 & 936 \\ 522 & 392 \\ 804 & 606 \\ 1504 & 1302 \\ 4.67 , df = 17 , p < 0.01 ; l^2 = 9 \\ 769 & 515 \end{array}$	99.4 %)	$\begin{array}{c} 0.406\ [0.401, 0.414\\ 0.430\ [0.382, 0.478\\ 0.432\ [0.382, 0.478\\ 0.432\ [0.382, 0.478\\ 0.479\ [0.413, 0.546\\ 0.502\ [0.458, 0.545\\ 0.510\ [0.482, 0.539\\ 0.566\ [0.551, 0.579\\ 0.575\ [0.573, 0.549\\ 0.660\ [0.524, 0.549\\ 0.660\ [0.524, 0.549\\ 0.566\ [0.561\ [0.570\ 0.576\\ 0.570\ [0.573\ 0.575\\ 0.751\ [0.713, 0.767\\ 0.754\ [0.733, 0.783\\ 0.666\ [0.493, 0.640\\ 0.667\ [0.493, 0.640\\ \end{array}$

Fig. 3 Forest plot for HPV vaccine acceptance rates among parents by world regions showing individual study estimates as well as pooled estimates (random-effects models) (n=62). CI=Confidence interval; df=Degrees of freedom; Events=The number of parents/guardians in each study who reported acceptance of the HPV vaccine for their children; $l^2 = I$ -squared statistic, indicating the percentage of variation due to heterogeneity; p < 0.01 = p-value, indicating statistical significance; Q=Cochran's Q statistic for heterogeneity; RE Model=Random-effects model; Total=The total number of parents/guardians in each study included in the meta-analysis

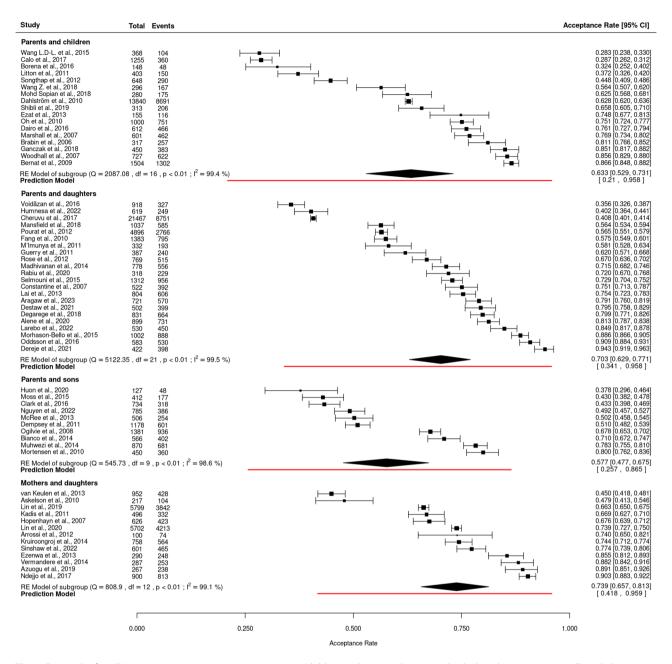


Fig. 4 Forest plot for HPV vaccine acceptance rates among parent/children's subgroups showing individual study estimates as well pooled estimates (random-effects models) (n = 62). CI = Confidence interval; df = Degrees of freedom; Events = The number of parents/guardians in each study who reported acceptance of the HPV vaccine for their children; $l^2 = I$ -squared statistic, indicating the percentage of variation due to heterogeneity; p < 0.01 = p-value, indicating statistical significance; Q=Cochran's Q statistic for heterogeneity; RE Model=Random-effects model; Total=The total number of parents/guardians in each study included in the meta-analysis

24.7 to 91.4% of parents [49–51]. In studies from North America (USA; n=1) and Oceania (New Zealand; n=1) the proportions were higher, with 95.8% and 92.6% of parents, respectively, recognizing this benefit [52, 53]. In contrast, South America (Brazil; n=1) had the lowest reported proportion, with only 10.0% of parents acknowledging cancer protection as a benefit [54].

Benefit 2: Recommendation by a pediatrician or family physician

Recommendations by a paediatrician or family physician were most frequently reported as a benefit in studies from North America (USA; n=3), where the proportion ranged from 22.0 to 77.0%, indicating strong influence from healthcare providers [52, 55, 56]. In Asia (India; n=1), a high proportion of 81.2% of parents considered healthcare recommendations a benefit, whereas in

Table 2 Descriptive analysis of seven parental barriers and benefits to HPV vaccinations reported for 43 studies

Africa (n = 12) Asia (n = 6) % (reference) % (reference)		= 6)	Australia		Europe (n = 11) % (reference)		North America (n = 10) % (reference)		Oceania (n = 1) % (reference)		South America (n = 2)		
		% (reference)		% (reference)							% (refe	erence)	
Benefi	1: Desire to protect their childr	en again	st cancer										
40.4%	Humnesa et al. 2021 ♀	91.4%	Madhivanan et al. 2014 Q	-		6.0%	Dahlström et al. 2010 ♀♂	95.8%	Guerry et al. 2011 ♀	92.6%	Rose et al. 2012 Q	10.0%	Mendes Lobão et al. 2018 🖓
77.0%	Sinshaw et al. 2022 ♀♀	82.3%	Degarege et al. 2018 ♀			67.0%	Mortensen et al. 2010 👌						
		24.7%	Alaamri et al. 2023 Q			45.7%	Bianco et al. 2014 3						
Benefi	2: Recommendation by a pedia	atrician o	r family physician										
-	-	81.2%	Degarege et al. 2018 ♀	-	-	17.9%	Bianco et al. 2014 🖑	22.0%	Reiter et al. 2009 ♀	-	-		
								75.0%	Guerry et al. 2011 ♀				
								77.0%	VanWormer et al. 2017 ♀♂				
Barrier	1: Concerned about vaccine ef	fectivene	955										
2.5%	Ezenwa et al. 2013 ♀♀	75.5%	Madhivanan et al. 2014 ♀	5.0%	Marshall et al. 2007 ♀♂	58.2%	Brabin et al. 2006 ♀♂	13.0%	Guerry et al. 2011 ♀	-	-		
25.6%	Sinshaw et al. 2022 ♀♀	81.0%	Degarege et al. 2018 ♀			79.4%	Dahlström et al. 2010 ♀♂	15.0%	Litton et al. 2011 ^a ♀♂				
21.1%	Aragaw et al. 2023 ♀♂					31.6%	Bianco et al. 2014 👌	0.3%	Hirth et al. 2019⁵ ⊊ੈ				
						8.1%	Borena et al. 2016 ♀♂						
						49.2%	Della Polla et al. 2020 ♀♂						
Barrier	2: Concerned about the advers	e effects	of the HPV vaccine										
\$1.7%	Vermandere et al. 2014 ♀♀	0.8%	Oh et al. 2010 ^c ♀	66.4%	Marshall et al. 2007 ♀♂	91.6%	Dahlström et al. 2010 ♀♂	9.5%	Litton et al. 2011ª ♀♂	-	-	61.0%	Mendes Lobão et al. 2018 Q
6.0%	Morhason-Bello et al. 2015 ♀♂	5.2%	Oh et al. 2010 ^c ්			87.5%	Gefenaite et al. 2012 ♀♀	39.0%	O'Leary et al. 2018 ♀				
25.6%	Dairo et al. 2016 ♀♂	28.9%	Songthap et al. 2012 ♀♂			30.2%	Borena et al. 2016 ♀♂						
3.0%	Azuogu et al. 2019 ♀♀	31.6%	Madhivanan et al. 2014 ♀			26.5%	Voidăzan et al. 2016 ♀						
17.8%	Dereje et al. 2021 ♀	76.0%	Degarege et al. 2018 Q			43.1%	Ganczak et al. 2018 ♀♂						
50.6%	Humnesa et al. 2021 ♀	56.9%	Shibli et al. 2019 ♀♂			41.8%	Della Polla et al. 2020 ♀♂						
1.9%	Larebo et al. 2022 ♀	18.8%	Alaamri et al. 2023 🎗										
7.6%	Mihretie et al. 2022 ♀♂												
52.4%	Sinshaw et al. 2022 ♀♀												
72.9%	Aragaw et al. 2023 ♀♂												
	3: Belief that vaccination will e	-											
6.4%	Ezenwa et al. 2013 ♀♀	54.6%	Songthap et al. 2012 ♀♂	4.9%	Marshall et al. 2007 ♀♂		Brabin et al. 2006 ♀♂	0.6%	Reiter et al. 2010 Q	8.8%	Rose et al. 2012 Q	-	
9.9%	Morhason-Bello et al. 2015 ♀♂	18.5%	Madhivanan et al. 2014 ♀			42.0%	Woodhall et al. 2007 ♀♂	34.7%	Guerry et al. 2011 ♀				
45.3%	Sinshaw et al. 2022 ♀♀	83.4%	Degarege et al. 2018 ♀			21.1%	Gefenaite et al. 2012 ♀♀	0.4%	Hirth et al. 2019⁵♀♂				
5.1%	Aragaw et al. 2023 ♀♂					20.6%	Bianco et al. 2014 ♂						
						11.0%	Oddsson et al. 2016 ♀						
Demis	4: Lack of recommendations					34.3%	Ganczak et al. 2018 ♀♂						
56.8%	Aragaw et al. 2023 ⊈∛	-	-	-	-	14.0%	Mortensen et al. 2010 ♂	8.8%	Fang et al. 2010 ♀	-	-	16.1%	Arrossi et al. 2012 ♀♀
						5.4%	Borena et al. 2016 ♀♂	59.6%	Guerry et al. 2011 ♀				
								14.0%	Gilkey et al. 2012ª ♀				
								27.0% 16.2%	Gilkey et al. 2012³♂ Hirth et al. 2019⁵♀♂				
Barrie	5: Lack of knowledge							10.270	1111111111112010 ‡0				
	Ezenwa et al. 2013 QQ	73.8%	Madhivanan et al. 2014 ♀			70.0%	Mortensen et al. 2010 ්	22.0%	Gottlieb et al. 2009 ♀				
13.9%	Vermandere et al. 2014 ♀♀	10.0%	Alaamri et al. 2023 ♀				Borena et al. 2016 ♀♂	65.5%	Guerry et al. 2009 ⊊			-	
3.7%	Morhason-Bello et al. 2015 ♀♂	10.070				/0		15.6%	Hirth et al. 2019 ^b ⊋∂				
42.2%	Dairo et al. 2016 ♀♂												
	Alene et al. 2020 ♀												
44.8%													

Q = Parents and daughters

 σ = Parents and sons

 $Q_{\mathcal{C}} = Parents and children$

QQ = Mothers and daughters

^a Mentioned by intenders

^b Mentioned by non-intenders

^c Oh et al. 2010 and Gilkey et al. 2012 provided valid percentage values for parents of daughters and parents of sons

Europe (Italy; n=1), only 17.9% of parents shared this view [47, 50].

Barrier 1: Concerned about vaccine effectiveness

Parental concern about vaccine effectiveness varied significantly in European studies (England, Italy, Poland, Romania, Sweden; n=5). Most studies (England, Italy, Sweden; n=4) reported moderate rates, ranging from 31.6 to 79.4% [46, 47, 57, 58]. In contrast, studies from North America (USA; n=3) and Africa (Ethiopia, Nigeria; n=3) showed generally low concern, with rates from 0.3 to 15.0% in North America [52, 59, 60] and 2.5 to 25.6% in Africa [45, 61, 62]. In Asian studies (India; n=2), the concern was much higher, ranging from 75.5 to 81.0% [50, 51]. In Australia (n=1), the concern was very low, with only 5.0% of the parents expressing doubts about vaccine efficacy [63].

Barrier 2: Concerned about the adverse effects of the HPV vaccine

Concern about the adverse effects of the HPV vaccine was most frequently reported in studies from Africa (Ethiopia, Kenya, Nigeria; n=10), with a wide range of perceptions: six studies indicated lower levels of concern, ranging from 1.9 to 25.6% [28, 64–68], while four studies reported moderate levels, ranging from 41.7 to 72.9% [44,

45, 62, 69]. In Asia (India, Israel, Korea, Saudi Arabia, Thailand; n=6), perceptions also varied, with three studies reporting low concern, ranging from 0.8 to 28.9% [49, 70, 71], and three indicating moderate concern, ranging from 31.6 to 76.0% [50, 51, 72]. In studies from Europe (Austria, Italy, Netherlands, Poland, Romania, Sweden; n=6), parents reported the highest perceived concern on adverse effects, with proportions ranging from 26.5 to 91.6% [25, 46, 57, 73–75]. In North America (USA; n=2), the concern ranged from 9.5 to 39% [60, 76]. In Australia (n=1) and South America (Brazil; n=1), the perceived concern were moderate, at 66.4% and 61.0%, respectively [54, 63].

Barrier 3: Belief that vaccination will encourage sexual activity (will lead to promiscuity)

The belief that HPV vaccinations might lead to unprotected sex was most prevalent in European studies (England, Finland, Iceland, Italy, Netherlands, Poland; n=6), with proportions ranging from 11.0 to 42.0% [25, 47, 58, 75, 77, 78]. The concern varied more widely in North American studies (USA; n=3), from 0.4 to 34.7% [52, 59, 79]. This belief was generally less common in studies from Africa (Ethiopia, 371 Nigeria; n=4), with three studies reporting proportions between 6.4 and 9.9% [61, 62, 66]. In Asian studies (India, Thailand; n=3), concern was more pronounced, ranging from 18.5 to 83.4% [50, 51, 71]. In contrast, Australia (n=1) and Oceania (New Zealand; n=1) reported the lowest concern, with 4.9% and 8.8% respectively [53, 63].

Barrier 4: Lack of recommendations

Lack of recommendations was most frequently reported as a parental barrier in studies from North America (USA; n=4), where the proportions ranged from 8.8 to 59.6%, indicating significant variation in its influence on vaccination decisions [52, 59, 80, 81]. In European studies (Austria, Denmark; n=2), the range was narrower and lower, from 5.4 to 14.0%, suggesting a more consistent but still notable barrier [48, 74]. In Africa (Ethiopia; n=1), a moderate proportion of 56.8% of parents reported lack of recommendations as a significant barrier, whereas in South America (Argentina; n=1), the proportion was lower at 16.1% [62, 82].

Barrier 5: Lack of knowledge

Lack of knowledge was most frequently reported as a parental barrier in studies from Africa (Ethiopia, Kenya, Nigeria; n=6), with proportions ranging from 3.7 to 51.5% [61, 66–69, 83]. In North American studies (USA; n=3), 15.6 to 65.5% of parents reported insufficient knowledge as influencing their decision not to vaccinate their children [52, 59, 84]. In Asian studies (India, Saudi Arabia; n=2), this reason was cited by 10.0 to 73.8% of

parents [49, 51]. In studies from Europe (Austria, Denmark; n=2), 22.7 to 70.0% of parents decided against vaccination due to lack of knowledge [48, 74].

Parental knowledge of HPV and HPV vaccinations

Parental knowledge of HPV and HPV vaccination was assessed as the proportion of the responses to five key HPV-related questions from 52 studies conducted in Africa (n=15), Asia (n=11), Europe (n=10), North America (n=12), Oceania (n=2) and South America (n=2) (Table 3).

In total, 46,905 parents aged 18 to >70 years answered one or several HPV-related questions. Five studies received a study quality rating of "very good", 18 studies were "good", 28 studies were "satisfactory", and one study was "unsatisfactory".

The proportion of parental knowledge level across the 52 studies varied widely according to the questions asked (6.5 to 100%), with 19 studies demonstrating high knowledge levels (\geq 80%), 18 studies indicating moderate knowledge levels, and 15 studies showing low knowledge levels (\leq 30%).

Knowledge about HPV was most frequently assessed (n=36), with the highest proportions observed among parents of daughters in a study from North America and the lowest among parents and children in a European study. Following this, awareness that cervical cancer is related to HPV infection was also widely reported (n=25), with the highest proportion found among parents of daughters in a study from North America and the lowest in an Asian study among parents of sons.

Knowledge question 1: Aware of HPV

Parental awareness of HPV was highest in North American studies (Canada, USA; n=9), ranging from 59.7 to 100% [52, 80, 84–90], reflecting consistently high awareness across the region. In studies from Africa (Ethiopia, Kenya, Morocco, Nigeria; n=8), awareness varied widely from 10.5 to 88.1%, with higher levels in studies from 2020 onward [44, 61, 64, 65, 68, 83, 91, 92]. Asian studies (China, India, Malaysia, Saudi Arabia, Thailand, Vietnam; n=8) showed similar variation, with rates from 26.5 to 81.2% [49, 50, 71, 72, 93-96]. In European studies (Austria, England, Finland, Italy, Poland, Romania, Spain; *n*=8), the responses varied from 11.0% to 94.4 [25, 57, 58, 73, 74, 78, 97, 98]. In studies from Oceania (New Zealand, Republic of Fiji; n=2), awareness was 10.2 to 59.0% [53, 99], and in South America (Brazil; *n*=1), it was 55.0% [54].

Knowledge question 2: Aware of the HPV vaccine

Parental awareness of the HPV vaccine was highest in European studies (Italy, Poland, Spain; n=3), ranging from 61.4 to 92.1% [57, 97, 98], showing consistently

Table 3 Descriptive analysis of parental knowledge level to HPV vaccinations reported for 52 studies

Africa (n = 15) % (reference)		Asia (n = 11) ^ª		Australia (n = 0)	Europe (n = 10)		North America (n = 12)		Oceani	a (n = 2)	South America (n = 2)	
		% (refe	rence)	% (reference)	% (refe	rence)	% (refe	rence)	% (refe	rence)	% (refe	ence)
Knowl	edge question 1: Aware of HPV											
24.0%	M'lmunya et al. 2011 ♀	70.8%	Songthap et al. 2012 ♀♂		11.0%	Brabin et al. 2006 ♀♂	69.8%	Hopenhayn et al. 2007 Q	59.0%	Rose et al. 2012 ♀	55.0%	Mendes Lobão et al. 2018 🤤
27.9%	Ezenwa et al. 2013 ♀♀	54.2%	Ezat et al. 2013 ⊊∂		79.0%	Woodhall et al. 2007 ⊊♂	72.6%	Ogilvie et al. 2008 ♂	10.2%	La Vincente et al. 2015 Q		
15.6%	Selmouni et al. 2015 Q	78.8%	Degarege et al. 2018 ♀		85.4%	Borena et al. 2016 ♀♂	80.8%	Bernat et al. 2009 ♀♂				
10.5%	Dairo et al. 2016⊋♂	42.3%	Lin et al. 2019 ♀♀		85.8%	Voidăzan et al. 2016 ♀	78.0%	Gottlieb et al. 2009 ♀				
12.8%	Azuogu et al. 2019 ♀♀	81.2%	Shibli et al. 2019 ⊊∂		55.3%	Ganczak et al. 2018 ♀♂	88.0%	Allen et al. 2010 Q				
55.2%	Alene et al. 2020 ♀	42.0%	Lin et al. 2020 ♀♀		94.4%	Della Polla et al. 2020 ♀♂	59.7%	Fang et al. 2010 ♀				
52.8%	Humnesa et al. 2021 ♀	26.5%	Nguyen et al. 2022 ♀		90.7%	López et al. 2022 ♀♂	72.4%	Guerry et al. 2011 ♀				
88.1%	Larebo et al. 2022 Q	68.0%	Alaamri et al. 2023 ♀		74.2%	Sobierajski et al. 2023 ♀♂	81.0%	Sadigh et al. 2012 ♀♀				
							100%	Lai et al. 2013 ♀				
	edge question 2: Aware of the H											
19.7%	Ezenwa et al. 2013 ♀♀	51.0%	Ezat et al. 2013 ♀♂		73.6%	Della Polla et al. 2020 ♀♂	65.0%	Allen et al. 2010 Q	-		36.5%	Arrossi et al. 2012 ♀
60.9%	Muhwezi et al. 2014 Q	21.0%	Lin et al. 2019 ♀♀		92.1%	López et al. 2022 ⊊∛	59.8%	Fang et al. 2010 ♀			89.0%	Mendes Lobão et al. 2018
6.5%	Dairo et al. 2016 ♀♂	84.0%	Shibli et al. 2019 ⊊∂		61.4%	Sobierajski et al. 2023 ♀♂	53.1%	Guerry et al. 2011 ♀				
73.0%	Dereje et al. 2021 ♀	20.5%	Lin et al. 2020 ♀♀				63.0%	Pourat et al. 2012 Q				
54.6%	Humnesa et al. 2021 ♀						84.0%	Sadigh et al. 2012 ♀♀				
84.9%	Larebo et al. 2022 Q											
48.7%	Mihretie et al. 2022 ♀♂											
73.2%	Sinshaw et al. 2022 ♀♀											
	edge question 3: Aware that HP		-									
74.1%	Ezenwa et al. 2013 ♀♀	88.2%	Songthap et al. 2012 ♀♂		89.7%	Bianco et al. 2014 3		Ogilvie et al. 2008 ở	56.7%	Rose et al. 2012 ♀	92.0%	Mendes Lobão et al. 2018 ♀
79.4%	Muhwezi et al. 2014 🖓	59.7%	Kruiroongroj et al. 2014 ♀♀		31.3%	Ganczak et al. 2018 ⊊∛	81.0%	Lai et al. 2013 ♀				
35.9%	Rabiu et al. 2020 Q	32.8%	Mohd Sopian et al. 2018 ♀♂		84.2%	Della Polla et al. 2020 ♀♂						
9.0%	Dereje et al. 2021 ♀	71.0%	Lin et al. 2019 ♀♀		89.2%	López et al. 2022 ⊊∛						
50.4%	Humnesa et al. 2021 ♀	62.0%	Shibli et al. 2019 ⊊∂									
57.9%	Larebo et al. 2022 ♀	69.6%	Lin et al. 2020 ♀♀									
37.2%	Mihretie et al. 2022 ♀♂	18.3%	Nguyen et al. 2022 ♀									
33.6%	Sinshaw et al. 2022 ♀♀											
Know	ledge question 4: Aware that ce	rvical can	cer is related to HPV infection									
11.0%		19.0%		· ·	23.8%	Borena et al. 2016 ♀♂	43.0%	Litton et al. 2011⁰ ⊋∂	51.4%	Rose et al. 2012 ♀	86.0%	Mendes Lobão et al. 2018♀
63.9%	Dairo et al. 2016 ♀♂	7.5%	Oh et al. 2010 ^ª ♂		30.5%	Oddsson et al. 2016 Q	95.0%	Lai et al. 2013 ♀				
11.0%	Dereje et al. 2021 ♀	87.3%	Songthap et al. 2012 ♀♂		36.2%	Ganczak et al. 2018 ⊊∛						
50.7%	Humnesa et al. 2021 ♀	55.5%	Ezat et al. 2013 ♀♂		73.7%	López et al. 2022 ♀♂						
73.0%	Larebo et al. 2022 Q	49.2%	Kruiroongroj et al. 2014 ♀♀									
32.1%	Mihretie et al. 2022 ♀♂	85.0%	Mohd Sopian et al. 2018 우강									
29.8%	Sinshaw et al. 2022 ♀♀	55.1%	Lin et al. 2019 ♀♀									
		73.8%	Shibli et al. 2019 ♀♂									
		53.1%	Lin et al. 2020 ♀♀									
		19.9%	Nguyen et al. 2022 ♀									
		62.3%	Alaamri et al. 2023 ♀									
Know	ledge question 5: Aware that the	e HPV vace	cine prevents cervical cancer									
81.5%		31.8%			71.4%	Bianco et al. 2014 👌		Lai et al. 2013 ♀	66.6%	Rose et al. 2012 ♀	65.0%	Mendes Lobão et al. 2018
60.9%	Muhwezi et al. 2014 ♀	57.5%	Lin et al. 2019 ♀♀				75.4%	O'Leary et al. 2018 ♀				
92.4%	Morhason-Bello et al. 2015 ♀♂		Shibli et al. 2019 ♀♂									
93.6%	Ndejjo et al. 2017 ♀♀	56.2%	Lin et al. 2020 ♀♀									
23.6%	Rabiu et al. 2020 ♀	65.2%	Alaamri et al. 2023 Q									
84.4%												
44.7%	Humnesa et al. 2021 ♀											
81.9%	Larebo et al. 2022 ♀											
59.4%	Mihretie et al. 2022 ♀♂											
71.4%	Sinshaw et al. 2022 ♀♀											

Q = Parents and daughters

d = Parents and sons

Q = Parents and children

QQ = Mothers and daughters

^a Oh et al. 2010 provided valid percentage values for parents of daughters and parents of sons

^b Mentioned by intenders

strong knowledge across the region. North American studies (USA, n=5) also has high awareness, between 53.1 and 84.0% [52, 80, 88, 89, 100]. South American studies (Argentina, Brazil; n=2) showed more variation, with levels from 36.5 to 89.0% [54, 82]. In contrast, studies from Africa (Ethiopia, Nigeria, Uganda; n=8) exhibits significant variation, with awareness ranging from 6.5 to 84.9% [28, 44, 45, 61, 64, 67, 68, 101], indicating lower knowledge in some regions. Asian studies (China, Israel, Malaysia; n=4) also varied widely, with awareness ranging from 20.5 to 84.0% [72, 93–95]. This highlights

greater variability in awareness in Africa and Asia compared to the consistently high levels in Europe and North America.

Knowledge question 3: Aware that HPV is sexually transmitted

Parental awareness that HPV is sexually transmitted was most consistent in North American studies (Canada, USA; n=2), with levels ranging from 74.2 to 81.0% [86, 90]. European studies (Italy, Poland, Spain; n=4) showed a broader range of awareness, from 31.3 to 89.7%, indicating varying levels of knowledge across the region [25, 47, 57, 97]. In studies from Asia (China, Israel, Malaysia, Thailand, Vietnam; n=7), awareness varies significantly, from low to high, with levels between 18.3 and 88.2%, highlighting considerable differences across countries [71, 72, 94–96, 102, 103]. African studies (Ethiopia, Nigeria, Uganda; n=8) had the widest range, from 9.0 to 79.4%, showing substantial variability in understanding across the region [28, 44, 45, 61, 64, 67, 101, 104]. Oceania (New Zealand; n=1) reports moderate awareness at 56.7% [53], while South America (Brazil; n=1) demonstrates high awareness at 92.0% [54].

Knowledge question 4: Aware that cervical cancer is related to HPV infection

Parental awareness of the link between HPV and cervical cancer is highest and most consistent in studies from North America (USA; n=2), ranging from 43.0 to 95.0% [60, 90]. European studies (Austria, Iceland, Poland, Spain; n=4) showed a range of awareness from low to moderate, spanning from 23.8 to 73.7% [25, 74, 77, 97]. Asian studies (China, Israel, Korea, Malaysia, Saudi Arabia, Thailand, Vietnam; n=10) exhibited the widest variability, with awareness ranging from 7.5 to 87.3% [49, 70–72, 93–96, 102, 103]. Similarly, studies from Africa (Ethiopia, Kenya, Nigeria; n=7) showed variability, with ranges from 11.0 to 73.0% [28, 44, 45, 64, 67, 68, 91]. Oceania (New Zealand; n=1) reports moderate awareness at 51.4% [53], and South America (Brazil; n=1) shows high awareness at 86.0% [54].

Knowledge question 5: Aware that the HPV vaccine prevents cervical cancer

In African studies (Ethiopia, Nigeria, Uganda; n=10), parental awareness that HPV vaccines prevent cervical cancer ranged from 23.6 to 93.6%, showing significant regional differences [28, 44, 45, 61, 64, 66, 67, 101, 104, 105]. In studies from Asia (China, Israel, Malaysia, Saudi Arabia; n=5), awareness was more consistent, ranging from 31.8 to 75.4% [49, 72, 94, 95, 103]. North American studies (USA; n=2) showed higher awareness levels at 75.4 to 93.0% [76, 90]. Europe (n=1; Italy), Oceania (n=1; New Zealand), and South America (n=1; Brazil) had similar levels of 71.4%, 66.6%, and 65.0%, respectively [47, 53, 54].

Discussion

This systematic literature review and meta-analysis summarized the global evidence on parental acceptance, attitudes, and knowledge regarding HPV vaccinations.

Parental acceptance of HPV vaccinations

Parents in the 86 included studies, totaling 251,880, generally supported HPV vaccinations for their children; Page 15 of 22

however, the HPV vaccine acceptance showed a high variation (12 to 97.5%). The acceptance of HPV vaccinations among parents was higher for daughters than for sons, and mothers were more likely to get their daughters vaccinated. This might be explained by insufficient knowledge of the parents on the vaccine's benefits for male children and adolescents and a perception of the vaccine as a preventative measure against cervical cancer. Sociocultural norms and targeted public health campaigns may also play a role.

The parents who were asked about their intention to vaccinate their children had higher acceptance rates than those whose children had received at least one dose of the HPV vaccine. No study characteristic that could explain the variation in the acceptance rates was identified. Higher acceptance rates in intention-based studies may reflect a general willingness but highlight barriers in translating intent into action, such as logistical challenges and lack of recommendations from healthcare providers.

The acceptance rates for HPV vaccinations in the cohort studies were significantly lower (5.0 to 36.8%) compared to the broader acceptance proportions reported in cross-sectional studies. The lower acceptance rates in these cohort studies, particularly in those that applied the Precaution Adoption Process Model (PAPM), were attributed to their specific focus on parents who had knowledge about the HPV vaccine but still remained hesitant [106, 107].

The meta-analysis showed that acceptance rates varied widely across the 62 studies (28.3 to 94.3%) over the years. The subgroup analysis revealed geographical variations in pooled parental HPV vaccine acceptance rates, with the highest rate observed in Africa and the lowest in North America. This is consistent with the results of other studies, which reported high parental acceptance rates in sub-Saharan Africa because of the available HPV vaccination programs and delivery vaccination strategies [108, 109]. In America, a moderate level of parental intent to have their children vaccinated against HPV was observed. Higher vaccination rates were reported when a healthcare provider recommended HPV vaccination to parents [110–114].

The geographical differences could be explained, potentially due to cultural and health political aspects, on the acceptance rates. Health policies, such as publicly funded national HPV vaccination programs, may enhance HPV acceptance rates. Nonetheless, the high HPV vaccine acceptance rates reported in some studies conducted in Africa or Asia are in contrast to the absence of such policies in these regions [115]. Another geographical factor could be cultural and socioeconomic aspects and attitudes toward vaccination in general. A recent study has demonstrated that low- and middle-income countries show a high acceptance rate of 80.0% for Covid-19 vaccines, whereas, in the USA, it was only 65.0% [116].

It is possible that these differences between low-/middle- and high-income countries are because individuals in high-income countries are more frequently exposed to public information regarding vaccines, safety concerns, side effects, and vaccine-related mortality. Additionally, inhabitants of countries with high availability and easy access to multiple vaccines may become less interested or be more skeptical toward vaccination in general because they may not see the need to receive yet another vaccination and are more exposed to information on vaccine authorization and safety [117].

The acceptance rate could also be correlated with the incidence of HPV in a given region, as the understanding and awareness of HPV could be higher if more individuals are infected. This assumption holds true for the comparison of America and Europe with comparatively lower HPV rates and Africa with the highest worldwide reported HPV rates [118]. Nonetheless, HPV infection rates in Asia are lower than in North America [119, 120], and therefore the high Asian acceptance rates cannot be explained by HPV incidence rates in this region. Moreover, the studies included in the present review do not allow for the conclusion that incidence rates in a given region correlate with acceptance rates, and hence, it remains to be determined to what extent these differences account for the different acceptance rates.

The perceived benefit of HPV vaccinations in preventing cancer and genital warts was the most frequently reported predictor of vaccine acceptance among parents. This is consistent with the findings of other systematic reviews, in which the perceived benefit in preventing cervical cancer not only was rated as the most important attribute of the HPV vaccine but also was positively associated with vaccine acceptance [21, 108]. These reviews had either a narrow population focus or a narrow geographic scope, and evidence on whether and how factors influencing parental HPV vaccine acceptance rates might differ across settings and populations is limited.

While most parents had a positive attitude toward HPV vaccinations, the fear of adverse effects was the most cited reason for vaccine non-acceptance, followed by doubts regarding the effectiveness and safety of the vaccine. This highlights the need for effective communication from health authorities and physicians to address these specific concerns and prevent parental misconceptions. Additionally, studies have confirmed that parents' belief that HPV vaccinations encourage unprotected sexual intercourse or early sexual activity decreases the likelihood that they will vaccinate their children [121–127]. According to prior research, concerns about the impact of the HPV vaccine on sexual behavior in children are unsubstantiated [128, 129]. Moreover, addressing

parents' concerns on the efficacy and safety of the vaccine could improve vaccination rates. There appears to be a large discrepancy between safety study results and parents' perception. Providing clear, evidence-based information and engaging healthcare providers in conversations with parents might help bridge this gap [130].

Parental attitudes regarding HPV and HPV vaccinations

Several studies have identified factors, such as the desire to protect their children against cancer, expectation that HPV vaccination is safe, and the level of knowledge associated with a parent's acceptance of HPV vaccines and their intention to vaccinate their children [23–25]. Support is generally high (74 to 78%) for a vaccine that protects against genital warts and cervical cancer but varies widely (12 to 100%), depending on the type of vaccine used and parents' ethnicity [131, 132].

In the US, the highest rates of cervical cancer have been observed among Latin-Americans or African-Americans; however, vaccination rates are suboptimal in these populations. For African-Americans, reasons for not getting their children vaccinated include concerns about the vaccine's efficacy and safety and their perception of a low infection risk, peer norms, and the perception of discrimination based on their socioeconomic status and their race [132–134]. For Latinos, vaccine acceptance depends primarily on the recommendations of their healthcare provider and the vaccine's ability to prevent cancer but also on fears of changes in the daughter's sexual behavior and her fertility as well as on individual and interpersonal reasons [132, 135, 136].

The initiation of HPV vaccinations largely depends on the parental intent to vaccinate their children, which is part of overall vaccine compliance. However, one study on HPV vaccine uptake among adolescents in the USA reported that factors that motivate parents to complete (or not complete) the vaccination series might differ, and the main reason for not intending to complete the HPV vaccine series once initiated was a lack of a recommendation from a healthcare provider for subsequent doses [112]. The parents of previously vaccinated children indicated that having their doctors or healthcare authorities recommend the vaccine increased their likelihood of having their child vaccinated against HPV. These results were consistent with those of previous studies in which practitioner recommendations appeared to be an important factor for vaccination acceptance [137–142]. The absence of a practitioner recommendation is an issue contributing to parental vaccine hesitancy. European studies have reported that nearly a third of hesitant parents never received any HPV vaccine recommendations [143]. It may be concluded that parents seem to trust the recommendations of their healthcare providers, and a lack of such recommendations most likely results in parents

hesitating to vaccinate their children and refusing HPV vaccinations.

Theoretical models have identified the intention to vaccinate as an important construct for vaccine-promoting interventions [23, 144]. According to the "Increasing vaccination model", vaccination rates may be increased by three interventions: modification of thoughts and feelings, motivation through social processes, and leveraging of people's thoughts and feelings [145]. Good knowledge regarding HPV and cervical cancer and positive attitudes toward vaccines have been associated with higher parental vaccine acceptance and intent to vaccinate [25, 146]. In contrast, barriers to HPV vaccination also affect the vaccine acceptance rates (defined as the willingness to be vaccinated) [135, 136]. Among parents, these barriers include the belief that their child is too young for HPV vaccination, vaccine-related safety concerns, and lack of knowledge [20-22, 110].

Parental knowledge of HPV and HPV vaccinations

The knowledge level regarding HPV infection and vaccinations varied widely within the population groups. In most studies, the knowledge level regarding HPV, that HPV is sexually transmitted, and HPV vaccinations provide protection against cervical cancer ranged from moderate to high. Several studies have confirmed that parents often express a need for more information regarding HPV vaccinations before deciding whether they should vaccinate their children [140, 147]. A systematic review has identified that insufficient knowledge levels (ranging from 67 to 81%) are a major barrier to HPV vaccinations among parents in Europe [148].

In several studies, parents have reported that they were unaware of the consequences of HPV or had never heard of HPV [60, 132, 149, 150]. However, other studies have indicated that HPV vaccine acceptance increases when parents are well informed about the benefits and risks of HPV vaccinations [24, 25]. A higher maternal level of education has been identified as a significant factor for increasing the acceptance and consequent uptake of HPV vaccinations [146, 151–156].

Limitations

A strength of this comprehensive review is that 86 studies covering an 18-year period (from 2006 to 2023) with predominantly large sample sizes were included. The main limitation is the considerable heterogeneity between the studies, including differences in the study design and reporting of ethnicity variables. Potential explanations for the residual heterogeneity include variations in the populations from whom the study samples were drawn, different vaccination delivery settings, and selection bias. Because of the differences in the measurements across the included studies and among the population groups, controlling the inconsistencies was difficult. Furthermore, while all eligible studies contained data on parental acceptance of HPV vaccination, not all studies included information on attitudes and knowledge. Although all studies were selected based on defined inclusion and exclusion criteria, the analysis was limited to the parameters that were reported, potentially affecting the comprehensiveness of our findings. We could not derive comprehensive findings on parental acceptance of HPV vaccinations for sons and the acceptance of HPV vaccinations by fathers separately. Overall, the frequent influence of small studies, high heterogeneity, wide PIs, and detected publication bias in certain subgroups underscores the need for cautious interpretation, particularly in cases where the evidence was rated as low or very low quality.

In summary, our findings highlight the importance of addressing the factors promoting HPV vaccine acceptance among parents and their children and the need for evidence-based interventions that address the widespread gaps between HPV vaccine recommendations and actual use [131]. A significant step in advancing acceptance is increasing parental knowledge on HPV, the effects of HPV infection, vaccine safety, and the availability of HPV vaccinations. Specifically targeting those groups of parents who may not be aware of the importance of vaccination in preventing cervical cancer and cancers affecting men, such as parents of sons, could enhance vaccine acceptance and vaccination rates in the future. Future vaccination policies should incorporate the WHO recommendations and consider that a high protection level is already achieved after one injection, which may be more achievable in many countries than a vaccination regimen encompassing two shots.

Conclusions

Parents' acceptance of HPV vaccines for their children was moderate in the studies included in this review, with notable variations across world regions. Acceptance rates were highest in Africa, likely due to effective public health campaigns, and lowest in North America, where safety concerns and logistical barriers were more prevalent. Some parents, particularly parents of sons, had limited knowledge on HPV infection and vaccinations, and they were less willing to vaccinate their sons compared to daughters. This suggests a need for targeted education and outreach efforts focused on the benefits of HPV vaccination for boys as well as girls. Given the body of evidence for the safety and effectiveness of HPV vaccinations, public knowledge about HPV infection should be promoted, including efforts to minimize the existing barriers to HPV vaccination and to increase vaccination accessibility and uptake. To counter misinformation and address safety concerns, national campaigns are required.

Healthcare providers may support this by informing parents about the safety and benefits of HPV vaccines. Improving parental attitudes toward HPV vaccinations is a key factor for increasing the moderate rates of parental acceptance of such vaccinations. Coordinated efforts among healthcare providers, parents, and government health authorities are urgently required to overcome the barriers to HPV vaccinations and increase parental acceptance.

Abbreviations

CI	Confidence interval
GRADE	Grading of Recommendations Assessment, Development, and
	Evaluation
HPV	Human papillomavirus
²	Inconsistency index
PI	Prediction interval
PRISMA	Preferred Reporting Items in Systematic Reviews and
	Meta-Analyses
PROSPERO	Prospective Register of Systematic Reviews
WHO	World Health Organization

Supplementary Information

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Supplementary Material 1	
Supplementary Material 2	
Supplementary Material 3	
Supplementary Material 4	

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Author contributions

SJK conceived and supervised the project. The lead author (SH) screened the titles and abstracts of the identified studies for inclusion. Two of the authors (SH, CC) independently reviewed the full texts of the studies and used standardized forms to extract data on the study characteristics. Two of the authors (SH, VO) independently assessed the quality of the included studies. SH and SJK conducted the meta-analyses and wrote the manuscript. GS reviewed and validated the results of the meta-analyses. All authors read and approved the final manuscript.

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Data availability

The datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

Declarations

Competing interests

The authors declare no competing interests.

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