



## REVIEW ARTICLE

# Epidemiological profile of human papillomavirus infections and cervical cancer prevention among sexually active women in Burkina Faso: Literature Review

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**ABSTRACT**

Cervical cancer is one of the biggest public health challenges in developing countries. According to the Global Cancer Observation in 2022, Burkina Faso recorded 988 new cases, with 775 deaths. Caused mainly by the human papillomavirus (HPV), this cancer is a heavy burden on our populations. The aim of this review was to assess the current state of human papillomavirus infections and cervical cancer prevention among sexually active women in Burkina Faso. Original studies were extracted from PubMed/Medline, Google Scholar, semanticscholar, Hinari and Science Direct using appropriate MeSH terms. Results were extracted and reviewed using the Preferred Reporting Items for Systematic Reviews and Meta-Analysis method (PRISMA). Twenty-four studies from three regions of Burkina Faso were included in the systematic review, with most of the articles from the "Centre" region. Out of a total of 5480 participants for the pooled analyses, the review showed a prevalence of HPV, varying according to population type and region and ranging from 20.6% to 87.2%. In contrast, the overall prevalence of HPV 16 and 18 infections were low compared to the global trend. With a low screening rate, the prevalence of cancerous lesions ranged from 1.5% to 15.42%. The level of knowledge was lower among rural than urban residents. Bivalent and quadrivalent vaccines had low coverage for the genotypes identified. In the fight against cervical cancer in Burkina, concerted efforts are needed to strengthen screening programs, increase HPV vaccination coverage, and raise public awareness. Large-scale national studies are needed to map HPV genotypes in order to make appropriate decisions for cervical cancer control.

**Keywords:** Epidemiology; Prevention; Cervical cancer; HPV; Burkina Faso.

## Abbreviations

HPV: Human Papillomavirus  
CC: Cervical Cancer  
GLOBOCAN: Global Cancer Observatory  
HR-HPV: High Risk Human Papillomavirus  
LR-HPV: Low risk Human Papillomavirus  
WHO: World Health Organization  
PCR: Polymerase chain reaction  
VIA/VILI: visual inspection with acetic acid or visual inspection with Lugol iodine  
HIV: Human Immunodeficiency Virus  
CIN: Cervical intraepithelial neoplasia  
KAP: Knowledge, Attitudes and Practices

## Introduction

Human Papillomavirus (HPV) is the world's most common sexually transmitted infection<sup>1</sup>. Today there are more than 200 HPV genotypes, with over 120 genotypes identified and sequenced<sup>2</sup>. Depending on their oncogenic potential, they are divided into two groups: high-risk oncogenic HPV (HR-HPV) and low-risk oncogenic HPV (LR-HPV). They are responsible for several infections of the epithelium of the anogenital tract and other mucous membranes, and are also involved in a variety of cancers, including vaginal, penile, vulvar, anal, and oropharyngeal cancers. Low Risk HPV such as HPV 6 and 11 are responsible for condylomatous lesions (warts) of the anogenital tract and laryngeal papillomatosis, most common benign tumor of the larynx<sup>3</sup>. Cervical cancer is an infectious pathology of the cervical mucosa<sup>4</sup>. In approximately 99% of cervical cancer cases, women have been exposed to HPV<sup>5</sup>. Several studies have shown that worldwide, HPV genotypes 16, 18, 31, 33, 35, 45, 52 and 58 are the most common among women with precancerous lesions and cervical cancer<sup>6-8</sup>. Some low-grade precancerous lesions induced by specific HPV genotypes have the potential to develop into cervical cancer if left untreated<sup>9</sup>. It is the fourth most common cancer in women worldwide, after breast, colorectal and lung cancers<sup>10</sup>. It is one of the world's major public health problems. In 2018, the World Health Organization (WHO) estimated that over 311000 women had died of cervical cancer and around 570000 new cases recorded worldwide. Around 85% of these deaths occurred in low- and middle-income countries<sup>11</sup>. Women in sub-Saharan Africa show the highest prevalence<sup>12</sup>, which appears to be rising steadily<sup>13</sup>.

According to the WHO, cervical cancer will kill more than 443,000 women a year worldwide by 2030, nearly 90% of them in sub-Saharan Africa<sup>13</sup>. According to GLOBOCAN, since 2020, cervical cancer has been the second leading cancer in terms of incidence and mortality in women in Burkina Faso, with 988 new cases and 775 deaths, as of 2022. For a more effective prevention of HPV infections, persistent HPV disease and cervical cancer, HPV vaccination is widely recommended<sup>14,15</sup>. The WHO estimates that HPV vaccination significantly

reduces cervical cancer morbidity and mortality. However, epidemiological data on HPV and cervical cancer from sub-Saharan African countries are necessary for the choice of available vaccines, as the distribution of HPV genotypes varies from one country to another<sup>9,14</sup>, with more genotype diversity in African countries<sup>16</sup>. The three licensed HPV prophylactic vaccines available are genotype-specific. In fact, the bivalent Gardasil (genotypes 16/18), the quadrivalent Cervarix (genotypes 6/11/16/18) and the nonavalent Gardasil 9 (6/11/16/18/31/33/45/52/58) have been shown to be safe and effective for primary prevention of the targeted genotypes<sup>17</sup>.

In Burkina Faso, since April 26, 2022, the quadrivalent human papillomavirus vaccine has been included in the vaccination schedule of the Expanded Program on Immunization to combat cervical cancer. However, are the genotypes identified in the general female population, in populations at high risk of HPV infection, in high-grade precancerous lesions and invasive cervical cancers predominantly covered by the vaccine available in the vaccination program? A systematic review of the epidemiology of human papillomavirus infection and cervical cancer in sexually active women in Burkina Faso therefore seems necessary to make better decisions on the control of HPV infections and cervical cancer, which carries a heavy burden in Burkina Faso. The aim of this review was to examine the epidemiological profile of HPV and the prevention of cervical cancer in sexually active women in Burkina Faso.

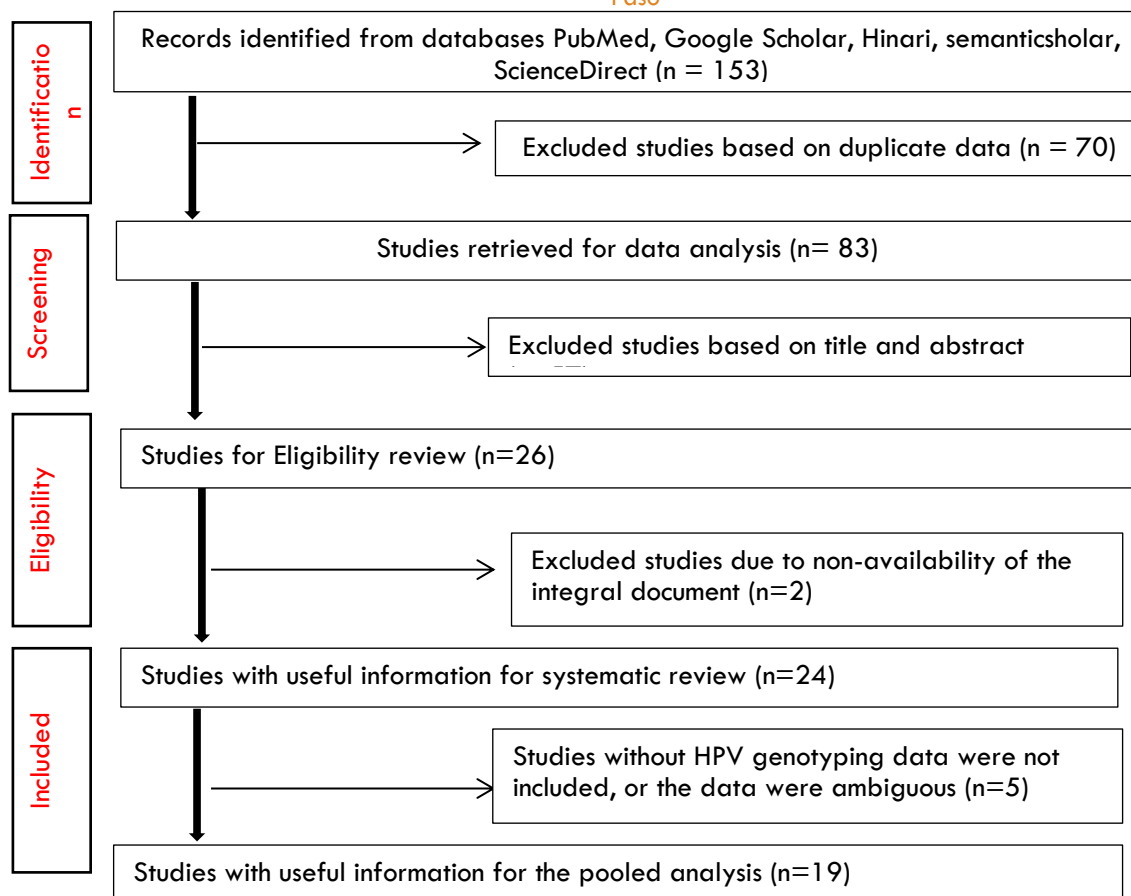
## Methods

### STUDY DESIGN

This study was conducted using the Preferred Reporting Items for Systematic Reviews and Meta-Analysis method (PRISMA)<sup>18</sup>. Research results and the number of articles included and excluded are presented in Figure 1.

### INFORMATION SOURCES AND RESEARCH STRATEGIES

We conducted a systematic literature review to identify relevant publications reporting on the epidemiology of human papillomavirus (HPV) infections and cervical cancer, as well as their prevention among sexually active women in Burkina Faso. Original studies in French and/or English were extracted from the PubMed/Medline, Google Scholar, semanticscholar, Hinari and Science Direct databases, using appropriate terms. The records identified have been downloaded in an appropriate format and linked to the zotero software. Boolean operators "AND" and "OR" were used to link mesh terms and to retrieve publications from PubMed/Medline (NCBI) databases. The keywords used were "Epidemiology OR prevention AND human papillomavirus OR cervical cancer" + "country name or country cities". The crude prevalence of HPV infection was calculated based on the crude data available in the eligible studies.



**Figure 1.** PRISMA diagram showing research strategy for the inclusion of published studies

#### ELIGIBILITY CRITERIA AND STUDY SELECTION PROCEDURE

After searching through the databases, the studies were then selected on the basis of the following criteria: data are published in a peer-reviewed scientific journal; articles are complete and available; articles relate only to patients residing in Burkina Faso; patients consulted for gynecological problems or participated in a cervical cancer screening campaign; cervical histology results are confirmed for patients with cervical cancer or precancerous lesions; HPV prevalence is calculated with at least five genotypes identified.

The articles selected for analysis will be used to determine the prevalence and genotypic distribution of oncogenic HPV in sexually active women in some of the regions of the country. The HPV molecular diagnostic techniques selected were those based on molecular biology techniques, in particular polymerase chain reaction (PCR) and hybrid capture. HPV genotype classification was also considered. In addition, we systematically excluded journal articles, correspondences to editors, press releases, letters, book chapters, publications in languages other than English/French, and studies whose data were ambiguous or could not be extracted. We considered HIV-negative and HIV-positive populations, sex workers, articles on the epidemiology and prevention of HPV and cervical cancer.

To get a true overview of circulating genotypes in Burkina Faso, HPV genotyping studies carried out on HPV-positive populations were selected. Studies on both Burkinabé and other female populations were included, but only data relating to the Burkinabé population were

considered in this study. Articles addressing other sexually transmitted pathologies in addition to HPV in the case of cervical cancer were included in this systematic review. However, articles from the Knowledge, Attitude and Practice surveys on health workers were excluded from this study. For pooled analyses, we considered the total number of HPV-diagnosed samples to be the size of our study population.

#### DATA EXTRACTION AND ANALYSIS

The data were extracted from various studies carried out in the regions of Burkina Faso for systematic examination. For the articles included in this review, the variables extracted were, study population and study setting, type of study or data collection (cross-sectional, prospective or retrospective) and method of detection. In addition, the overall crude prevalence of HPV infection was determined as the ratio of the total number of women tested positive for at least one HPV to the total number of samples tested, expressed as a percentage. The frequency of HPV genotypes was calculated as the ratio of the genotype of interest to the total number of genotypes identified, taking into account single, multiple and undetermined genotypes. IBM SPSS Statistic 26 and Excel 2016 were used for frequency calculations and graphing.

## Results

#### SELECTION OF STUDIES FOR REVIEW

In our study, a total of 153 articles were extracted from the databases, and for the systematic review, 26 full-text articles were included. According to our inclusion criteria, 24 scientific studies on epidemiological studies of HPV and cervical cancer prevention in sexually active women

conducted in 3 regions of Burkina Faso, including 19 articles on HPV genotyping and 5 articles on cervical

cancer screening, knowledge, attitudes, and practices, were selected (Table 1).

**Table 1.** Overview of original articles selected for the systematic review study

Population	Reference	Region (cities)	Area of interest	Average age (age range)	Size	HPV Prevalence (%)
SA	Kabre et al <sup>19</sup>	Centre (Ouagadougou)	HPV genotyping	28.2 (18 - 40)	100	23
HIV+/HIV-	Djigma et al <sup>20</sup>	Centre (Ouagadougou)	HPV genotyping	33.91 (20 - 53)	421	34.4
SA	Ouedraogo et al <sup>21</sup>	Centre (Ouagadougou)	HPV genotyping	31 (15 - 63)	300	43
SA	Salambanga et al <sup>22</sup>	Centre (Ouagadougou)	HPV genotyping	34.7 (18-57)	234	52.56
HIV-/HIV+	Djigma et al <sup>23</sup>	Centre (Ouagadougou)	HPV genotyping	34.1 (20 - 54)	250	59.6
SA/HIV+	Zohoncon et al <sup>24</sup>	Centre (Ouagadougou)	HPV genotyping	33.7 (20-53)	180	73.33
SA	Ouedraogo et al <sup>25</sup>	Centre-Est (Tenkodogo)	HPV genotyping	35.5 (20 - 60)	131	34.4
SA	Ouedraogo et al <sup>26</sup>	Centre-Est (Garango)	HPV genotyping	39.2	135	43
SA	Ouedraogo et al <sup>27</sup>	Centre, Centre-Est, Hauts-Bassins	HPV genotyping	28.02 (15 - 57) 37.37 (20 - 60) 33.14 (16 - 76)	520 266 535	35.42
SA	Sawadogo et al <sup>28</sup>	Centre (Ouagadougou)	KAP	29.37 (20-50)	840	-
SA	Compaore et al <sup>29</sup>	Centre (Ouagadougou)	CC screening & KAP	37.5 (18 - 72)	351	-
SA	Kagoné et al <sup>30</sup>	Hauts-Bassins (Bobo-Dioulasso, Bama)	CC screening & KAP	34.9 (18 - 60)	577	-
SA	Tassebedo et al <sup>31</sup>	Centre-Nord (Boussé)	CC screening & KAP	34 IQR (30-50)	418	-
SA	Diendéré et al <sup>32</sup>	National	CC screening & KAP	37.5 (25 - 49)	2293	-
SA	Traore et al <sup>33</sup>	Hauts-Bassins (Bobo-Dioulasso)	CC screening & HPV genotyping	35.3 (20-56)	181	25.4
SA	Ouattara et al <sup>34</sup>	Hauts-Bassins (Bobo-Dioulasso)	CC screening & HPV genotyping	30.7 (19 - 51)	234	20.6
SA/SW	Tovo et al <sup>35</sup>	Centre (Ouagadougou)	HPV genotyping	27.12 (17 - 50)	182	54.94
SA/SW	Ilboudo et al <sup>36</sup>	Centre (Ouagadougou)	HPV genotyping	27.3 (16 - 50)	200	53
SA/CC	Ouédraogo et al <sup>37</sup>	Centre (Ouagadougou)	HPV genotyping	41.5 (22 - 74)	43	48.8
SA/CC	Zohoncon et al <sup>38</sup>	Centre (Ouagadougou)	HPV genotyping	46.32 (21-84)	65	72.31
SA	Maria et al <sup>1</sup>	Centre (Ouagadougou)	HPV genotyping	41.3 (19 -76)	444	42.3
CC				42.3 (24 - 70)	39	87.2
SA/HBV/HCV	Ouédraogo et al <sup>39</sup>	Centre (Ouagadougou)	HPV genotyping	37.8 (20 - 65)	100	28
SA/HIV+	Chikandiwa et al <sup>40</sup>	Centre (Ouagadougou)	HPV genotyping	36 (IQR, 31-42)	594	27.1
SA/HIV+/SW	Didelot-Rousseau et al <sup>41</sup>	Hauts-Bassins (Bobo-Dioulasso)	CC screening & HPV genotyping	28 (16 - 54)	360	66.1
<b>Total</b>	<b>24 studies</b>	<b>3 Regions</b>	-	-	<b>9993</b>	<b>-</b>

**Legend:** SA: sexually active women; HIV+/- : HIV seropositive or seronegative; HCV: Hepatitis C virus; HBV: Hepatitis B virus; SW : sex workers; CC: Cervical Cancer; KAP: knowledge, attitudes and practices.

A total of 9993 participants from the sexually active female population were included for data extraction.

#### CHARACTERISTICS OF INCLUDED STUDIES

Studies included in this systematic review provided data from the Burkinabe population and met our selection criteria. Table 4 shows that among 5480 sexually active women in the general population, 2232 were infected with at least one of the HR-HPV (16/18/31/33/35/39/45/51/52/56/59/66/68/82) and/or LR-HPV (6/11/40/41/42/43/44/55/54/61/62/82/67/69/71/70/72/73/74/84) genotypes. Of the 24 scientific studies included in the review, 19 had information on the genotypic distribution of HPV and were therefore, eligible for pooled analysis. Multiple infections with at least two specific HPV types were reported in the majority of studies.

#### EPIDEMIOLOGY OF HUMAN PAPILLOMAVIRUS INFECTION

##### Prevalence of Human Papillomavirus infection in Burkina Faso

This systematic review reports on the prevalence and distribution of HPV genotypes among sexually active

women in Burkina Faso. The analysis of selected studies shows that the prevalence of HPV infections is very high in the general population of women in this low-income country.

HPV prevalence varied according to population type and locality in Burkina Faso. In the sexually active female population with unknown risk of HPV infection, it ranged from 20.6% to 52.56% in the studies included in this review.

Among the studies analyzed, 11 involved HIV-positive women, cases of high-grade precancerous lesions (CIN1/2/3) or cervical cancer, sex workers and HBV/HCV-positive women, thus representing populations at high risk of HPV infection. Indeed, these studies reported that the maximum prevalence of HPV in this at-risk population was 87.2%, 66.1%, 97.59%, 28% respectively in confirmed cases of precancerous lesions or cervical cancer, sex workers, HIV-positive women, and those with hepatitis B and/or C (Table 2).

**Table 2.** Overview of original articles on Human Papillomavirus infection in sexually active women

Study population	Reference	Region	Prevalence (HPV /Size)	HR-HPV	LR-HPV	Unspecified	Single genotype	Multiple genotypes	Nine most frequent HR-HPV genotypes, in descending order
<b>Population of women with unknown history of HPV infection</b>									
SA	Ouedraogo et al <sup>21</sup>	Centre	24.33 % (73/300)	53	20	45	64	9	50'S/18/16/30'S/45
SA	Salambanga et al <sup>22</sup>	Centre	52.56 % (123/234)	123	-	0	58	65	59/66/56/45/58/39/51/68/52
SA	Ouedraogo et al <sup>25</sup>	Centre-Est	34.4 % (45/131)	45	-	0	25	20	56/66/68/51/18/31/35/52/58
SA	Ouedraogo et al <sup>26</sup>	Centre-Est	43 % (58/135)	58	-	0	49	9	56/18/68/66/59/58/35/52/51
SA	Ouedraogo et al <sup>27</sup>	Centre Centre-Est Hauts-Bassins	35.42 % (468/1321)	225 103 140	- - -	0 0 0	116 74 108	109 29 32	56/52/66/59/39/51/18/35/68
SA	Traore et al <sup>33</sup>	Hauts-Bassins	25.4 % (46/181)	46	-	0	39	7	39/52/18/35/31/68/66/45/51
SA	Ouattara et al <sup>34</sup>	Hauts-Bassins	20.6 % (48/234)	48	-	0	39	9	52/66/68/39/51/18/31/35/56
SA	Kabre et al <sup>19</sup>	Centre	23 % (23/100)	16	13	2	12	9	18/35/31/52/58/56/45/59/33
SA	Maria et al <sup>1</sup>	Centre	42.3% (188/444)	127	108	0	96	92	16/52/18/35/59/66/68/39/53
<b>SA</b>	Djigma et al <sup>20</sup>	Centre	24.8% (59/238)	59	12	0	38	21	18/52/58/35/16/31/56
SA	Zohoncon et al <sup>24</sup>	Centre	30.2 (19/63)	19	-	0	9	10	52/18/31/35/45/51/39/56/58
<b>Total 1</b>			<b>34.01 (1150/3381)</b>	<b>1062</b>	<b>153</b>	<b>47</b>	<b>727</b>	<b>421</b>	<b>-</b>
<b>Population of women at high risk of HPV infection</b>									
HIV+	Djigma et al <sup>20</sup>	Centre	59.6 % (149/250)	124	25	119	102	47	18/50'S/30'S/16/45
SA/ HIV+	Djigma et al <sup>23</sup>	Centre	63.9 % (117/183)	117	6	0	44	73	18/35/31/52/58/56/45
SA/HIV+	Chikandiwa et al <sup>40</sup>	Centre	27.1 % (161/594)	-	161	0	127	34	Only low-risk HPV
SW	Didelot-Rousseau et al <sup>41</sup>	Hauts-Bassins	66.1 % (238 /360)	178	53	-	112	126	52/58/35/51/16/31/18/68/39
SW	Tovo et al <sup>35</sup>	Centre	54.94 % (100/182)	100	-	0	-	-	68/31/51/52/56/35/66/58/45
SW	Ilboudo et al <sup>36</sup>	Centre	53 % (106/200)	106	-	0	49	57	68/31/52/51/56/66/58/35/39
CU	Zohoncon et al <sup>38</sup>	Centre	72.30 % (47/65)	47	-	0	31	16	18/31/39/45/16/35/58/52/51
CIN2/3	Ouédraogo et al <sup>37</sup>	Centre	48.8 % (21/43)	21	-	0	19	2	39/35/45/33/51/52/56/18/31
HIV+	Zohoncon et al <sup>24</sup>	Centre	97.59% (81/83)	<b>81</b>	<b>-</b>	<b>0</b>	<b>8</b>	<b>73</b>	35/18/31/52/56/58/59/51/33/45/16
CIN	Maria et al <sup>1</sup>	Centre	87.2% (34/39)	30	5	0	14	20	52/18/35/59/66/68/39/53
SA/ VHB/ VHC	Ouédraogo et al <sup>39</sup>	Centre	28 % (28/100)	18	21	0	13	15	52/18/68/16/31/82/56/66/58
<b>Total 2</b>			<b>51.55%(1082/2099)</b>	<b>822</b>	<b>271</b>	<b>119</b>	<b>519</b>	<b>463</b>	<b>-</b>
<b>Total 1 and 2</b>			<b>40.73%(2232/5480)</b>	<b>1884</b>	<b>424</b>	<b>166</b>	<b>1246</b>	<b>884</b>	<b>18/52/56/35/31/58/66/59/51</b>

Legend: SA: sexually active women; HIV+/-: seropositive or seronegative HIV; SW: sex workers; CC: cervical cancer; CIN: Cervical intraepithelial neoplasia.



The pooled analysis reported that the overall prevalence of HPV in these different at-risk populations was 69.39%, 59.84%, 45.77%, 28% respectively in confirmed cases of precancerous lesions or cervical cancer, sex workers,

HIV-positive women, and HBV/HCV-positive women. This prevalence was 34.01% for the population with unknown high risk of HPV infection (Figure 2).

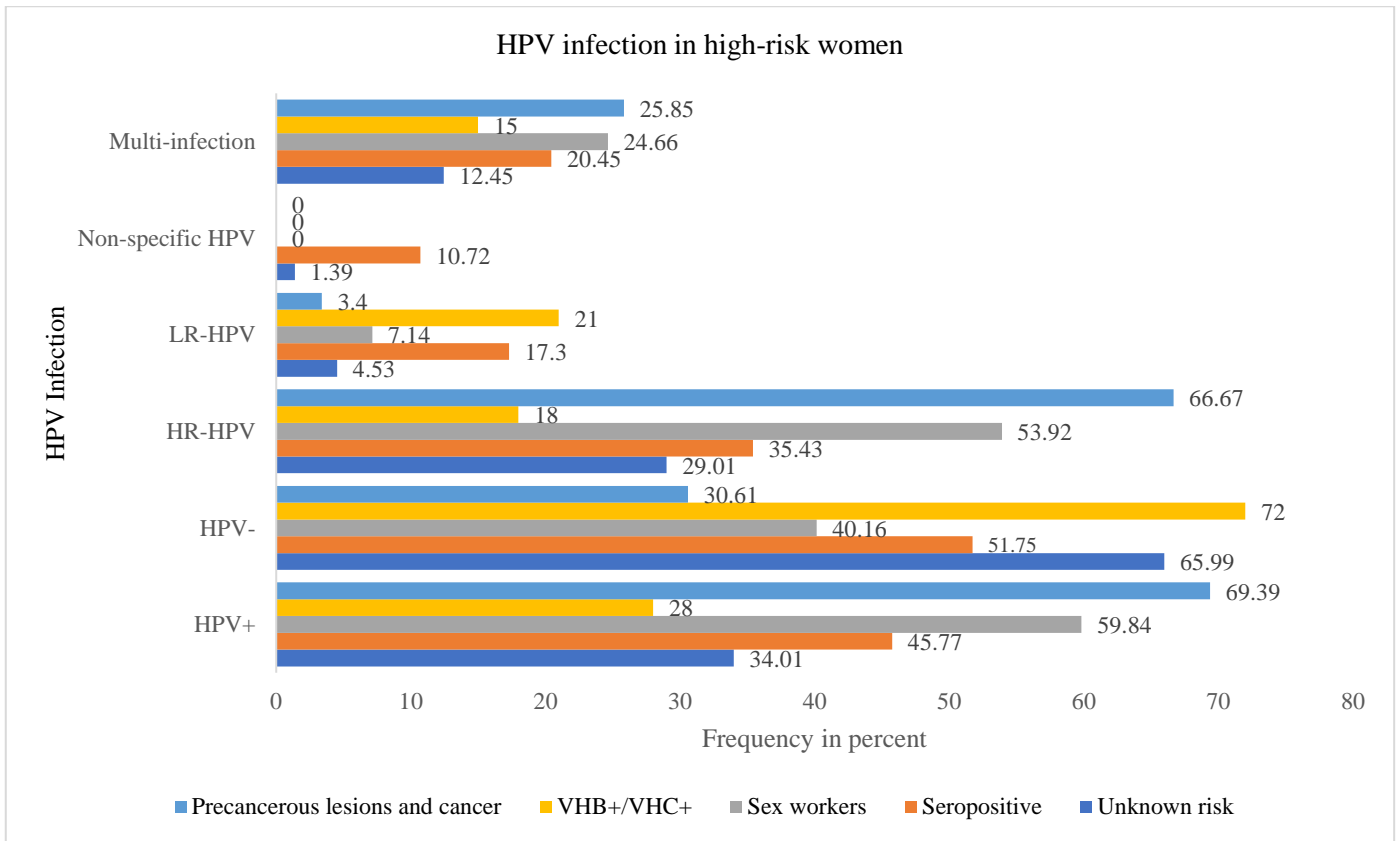


Figure 2. Human Papillomavirus infections in high-risk women

From the pooled analysis of data extracted from eligible studies, only three regions benefited from the HPV epidemiological studies, with the majority conducted in the “Centre” region. Indeed, among the 5480 participants, 3647 were from the “Centre” region, 1201 from the “Hauts-Bassins” region and 532 from the “Centre-Est” region. HPV prevalence reached 40.72% (2232/5480) in the general female population, 51.55%

(1082/2099) in the population at high risk of HPV infection, and 34.01% (1150/3381) in the female population with unknown history of HPV risk. In addition, among HPV-infected women, single and multiple infections were 55.82% (1246/2232) and 39.60% (884/2232) respectively, versus 7.44% (166/2232) of unspecified HPV infections.

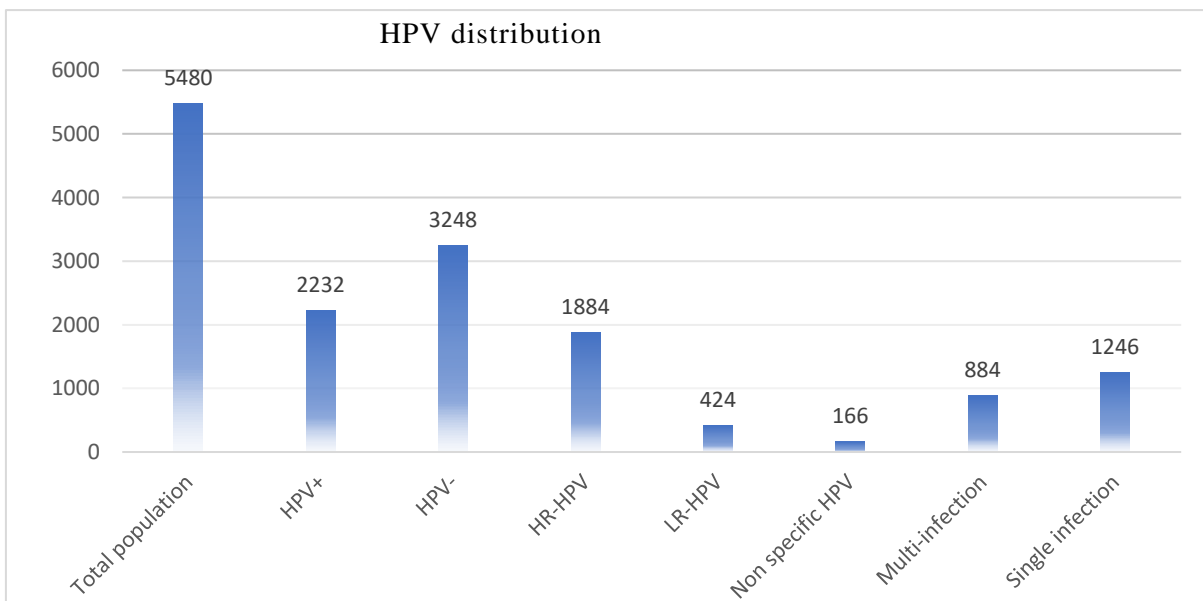


Figure 3. Overall population distribution by Human Papillomavirus status

Figure 4 shows the distribution of the female population according to HPV status by region.

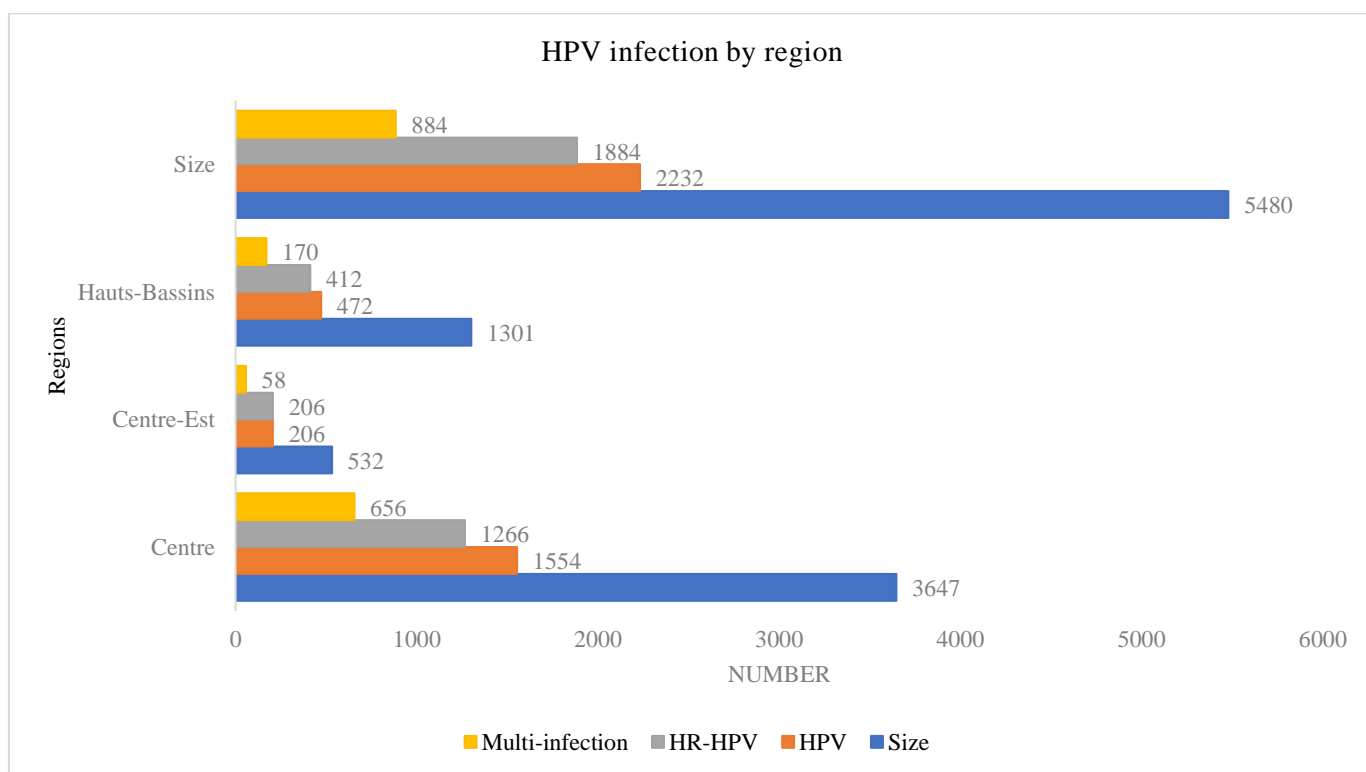


Figure 4. Distribution of women by HPV status and regions

### Human Papillomavirus genotype distribution

The studies included in this review showed that the predominance of HPV genotypes in women was a population-based and locality-based variable (Table 2). In fact, several high-risk and low-risk oncogenic HPV genotypes were found in the pooled analysis of 19 studies (Table 3). Among the genotypes, the most

frequent HR-HPV were HPV 16, 18, 31, 33, 35, 39, 45, 51, 52, 56, 58, 59, 66, 68 and the most frequent LR-HPV were HPV 6, 11, 43, 44/55, 54, 69/71, 70 (Table 3). Considering the three most predominant genotypes in the population at high risk of HPV infection, the studies show, in order of respective presence, HPV 18; 31; 52/35/68; 39; 45; 16 (Table 2).

Table 3. Prevalence of HPV genotypes in sexually active women based on collective analysis

HR-HPV	Number (%)	LR-HPV	Number (%)
HPV 16	153 (6.85)	HPV 6	101 (4.52)
HPV 18	<b>308 (13.80)</b>	HPV 11	11 (0.49)
HPV 31	<b>195 (8.74)</b>	HPV 40	10 (0.45)
HPV 33	74 (3.31)	HPV 42	4 (0.17)
HPV 35	<b>230 (10.30)</b>	HPV 43	17 (0.76)
HPV 39	153 (6.85)	HPV 44/55	56 (2.50)
HPV 41	2 (0.09)	HPV 54	29 (1.30)
HPV 45	146 (6.54)	HPV 61	3 (0.13)
HPV 51	<b>173 (7.75)</b>	HPV 62/81	14 (0.63)
HPV 52	<b>298 (13.35)</b>	HPV 67	11 (0.49)
HPV 53	36 (1.61)	HPV 69/71	25 (1.12)
HPV 56	<b>280 (12.54)</b>	HPV 70	26 (1.16)
HPV 58	<b>182 (8.15)</b>	HPV 72	4 (0.18)
HPV 59	<b>175 (7.84)</b>	HPV 74	37 (1.66)
HPV 66	<b>182 (8.15)</b>	HPV 84	1 (0.04)
HPV 68	134 (6.00)	<b>Total</b>	<b>349</b>
HPV 82	19 (0.85)		
<b>Total</b>	<b>2740</b>		

### Risk factors associated with Human Papillomavirus infections

Several risk factors favoring HPV infections have been described worldwide. In Burkina Faso, HIV/AIDS, age, multiparity, residence, early age at 1st sex, use of oral

contraception, condom use, level of education, number of sexual partners, etc., have been described as risk factors for HPV infection by the eligible studies included in this systematic review (Table 4).

**Table 4.** Overview of risk factors for HPV infection according to the studies selected

Article	Size	HPV infection risk factors (p-value)
Ouedraogo et al <sup>27</sup>	1321	- Level of education (0.002) - Marital status (0.001) - Occupation (< 0.001) - Number of sexual partner (0.003) - Frequency of intercourse (<0.001)
Salambanga et al <sup>22</sup>	234	- Age (0.037) - Sexual practices (0.001)
Ouédraogo et al <sup>39</sup>	100	- Education level (0.010) - Parity (0.006) - HIV status (0.002)
Maria et al <sup>1</sup>	410	- Place of birth (0.005) - Single (0.026) - Age at first sex < 20 years (0.001)
Kabre et al <sup>19</sup>	100	- Number of sexual partners (0.04)
Ouedraogo et al <sup>26</sup>	135	- Age at first sex (0.003) - Number of sexual partners (< 0.001) - Condom use (0.001)
Ilboudo et al <sup>36</sup>		- Level of education (0.019) - Number of sexual partners (< 0.001) - Condom use (< 0.001)

**EPIDEMIOLOGY OF CERVICAL CANCER**

**Prevalence, incidence, mortality and morbidity**

For more than a decade, cervical cancer has been a major public health problem in Burkina Faso, despite the efforts made to prevent it. According to GLOBOCAN,

since 2008 the crude incidence rate per 100,000 people has fluctuated between 15 and 26, and the crude morbidity rate per 100,000 people between 11 and 21 (Table 5).

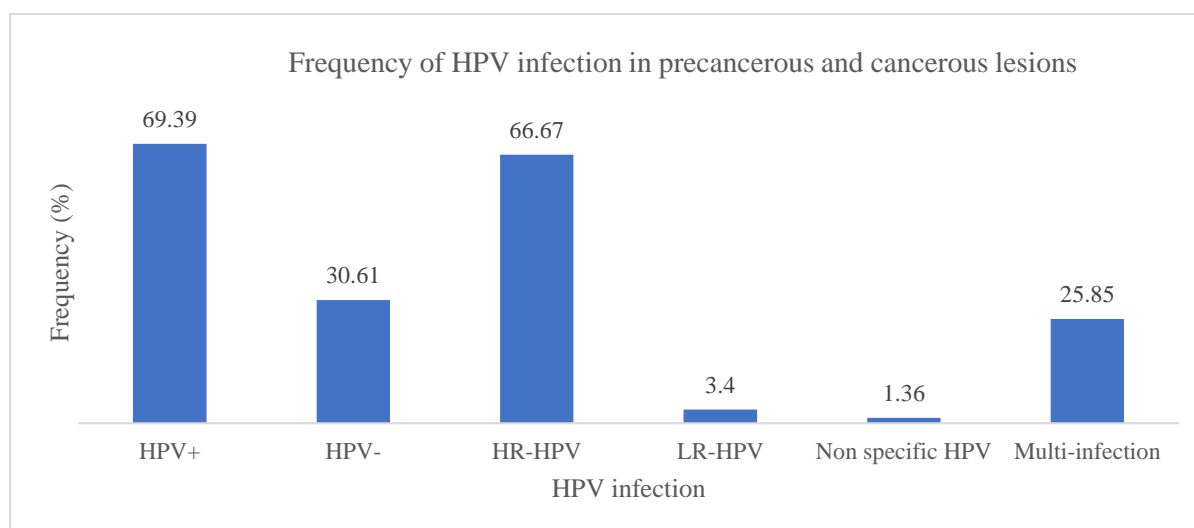
**Table 5.** Summary of GLOBOCAN incidence and morbidity rates over time

Year	Incidence		Morbidity	
	Number of cases	Crude rate/100000	No. of deaths	Crude rate/100000
2008	1230	16.1	838	11.0
2018	2517	25.4	2081	21.0
2020	1132		839	
2022	988	15.9	775	13.0

**Influence of Human Papillomavirus infection and Human Papillomavirus genotype distribution on the occurrence of cervical cancer in Burkina Faso**

According to the pooled analyses in this study, around 70% of women with precancerous lesions and cervical

cancer are infected with at least one HPV genotype, and around 67% are infected with at least one high-risk oncogenic HPV genotype (Figure 5).

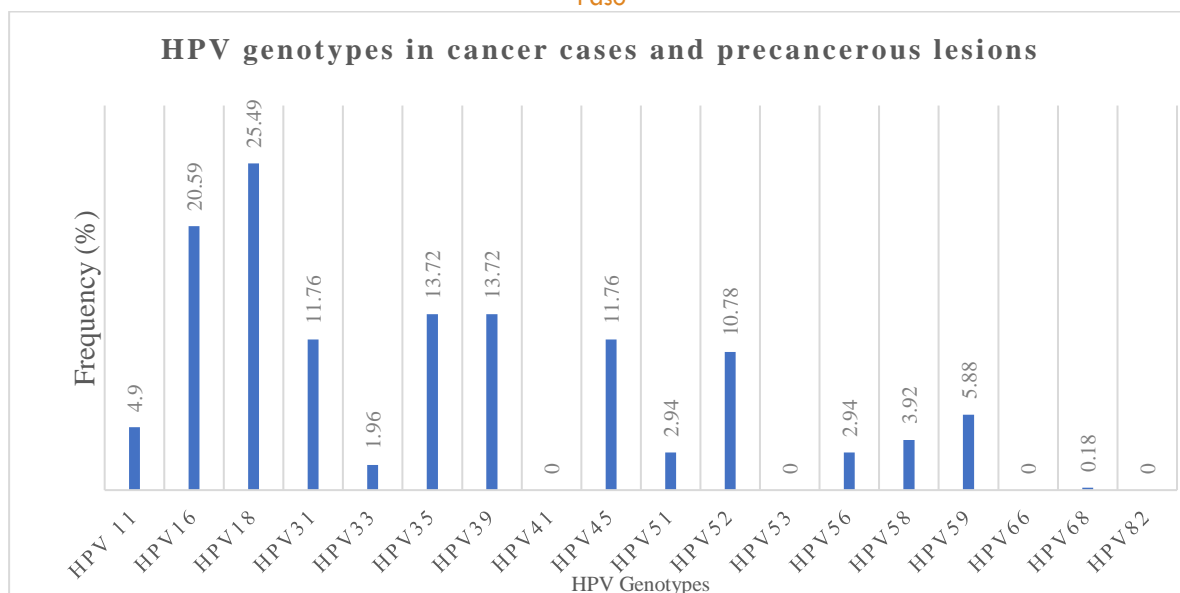


**Figure 5.** Frequency in percentage of Human Papillomavirus infection in precancerous lesions and cervical cancer

Among the high-risk oncogenic genotypes, HPV18, HPV16, HPV35 and HPV39 are the most frequent, with prevalence rates of 25.49%, 20.59% and 13.72% respectively (Figure 6). The predominant genotypes in

these cases of cancer, precancerous lesions and the population at high risk of infection in general, testify to the importance of a nonavalent vaccine that will include HPV-6/11/16/18/31/33/45/52/58.





**Figure 6.** Human Papillomavirus genotypes in cancer cases and precancerous lesions

**PREVENTION OF HUMAN PAPILLOMAVIRUS INFECTION AND CERVICAL CANCER**

**Risk factors associated with cervical cancer**

In addition to persistent infection with high-risk oncogenic HPV genotypes, considered to be the main cause of precancerous lesions and cervical cancer, several risk factors were associated. According to the articles included in this review, these were HIV serology, age, knowledge of risk factors and parity (Table 6).

**Screening and Knowledge, attitudes and practices on cervical cancer in Burkina Faso**

The knowledge, attitudes and practices of sexually active women towards Cervical cancer remain major prevention factors. In fact, the studies selected for this review revealed a low level of cervical cancer knowledge. Most women are still unaware of preventive measures, and less than 10% of women in the various targeted studies were screened.

Screening detects pre-cancerous lesions, which represent the sub-clinical or asymptomatic stage of the disease, and which, if untreated, can progress to cancer. Its aim is to identify individuals with a high probability of contracting or developing the disease. In low-income countries, screening by visual inspection with acetic acid and Lugol's iodine (VIA/VILI) is recommended by the WHO in addition to cervical smear. With this technique, the frequency of detection of precancerous lesions varies from 1.5% to 24% across the studies included. With a low screening rate as previously suggested, several factors such as level of education, age and residence were associated with screening, according to the studies examined. Like HPV infection, several factors are thought to be associated with the development of cervical cancer in women (Table 6).

**Table 6.** Screening and knowledge, attitudes and practices for cervical cancer in women in Burkina Faso

Article	Size	Risk factors associated with precancerous lesions and cervical cancer (p-value)		
Kagoné et al <sup>30</sup>	577	- Age (0.0005) - Parity (0.0442)		
Ouédraogo et al <sup>37</sup>	43	- HPV status (< 0.001)		
Didelot-Rousseau et al <sup>41</sup>	379	- Age (0.001) - HR-HPV types (0.002) - HIV-1 serology (< 0.001)		
Sawadogo et al <sup>28</sup>	840	- Think at risk of cervical cancer (0,02) - Family history of cervical cancer (0.002) - History of STI (0.01) - Heard about HPV (0.02)		
Screening and related factors				
Article	Size	Technic	Lesion frequency	Screening-related factors
Compaore et al <sup>29</sup>	351	-	-	-Education (<0.001) -Age (<0.001) -Residence (0.005) -Employment (0.004) -Motivation by a healthcare professional (0.041)
Kagoné et al <sup>30</sup>	577	VIA	15.42%	-
Tassebedo et al <sup>31</sup>	418	VIA/VILI	5%	-
Ouedraogo et al <sup>27</sup>	1321	VIA/VILI	4.39%	-
Didelot-Rousseau et al <sup>41</sup>	379	Cervical smear	24,0 %	-
Diendéré et al <sup>32</sup>	2293	-	-	- Geographic Regions (0.0001) - Residency (0.035)

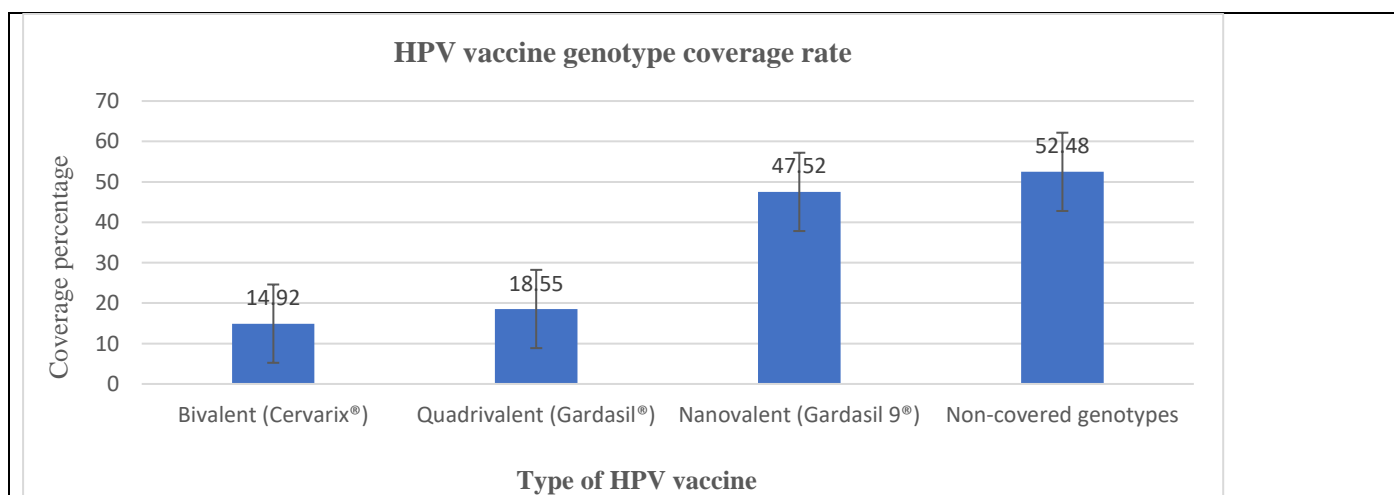
Article	Size	Risk factors associated with precancerous lesions and cervical cancer (p-value)		
				- Marital status (0.035) - Education (0.0001) - Occupation (0.0001)
Traore et al <sup>33</sup>	181	VIA/VILI	5% (9/179)	-
Ouattara et al <sup>34</sup>	234	VIA/VILI	7% (16/234)	-
Ouedraogo et al <sup>25</sup>	131	VIA/VILI	1.5% (2/131)	-
knowledge, attitudes and practices on cervical cancer				
Article	Size	knowledge, attitudes and practices (frequencies in %)		
Kagoné et al <sup>30</sup>	577	- Heard about cervical cancer (49%) - Knows at least one preventive measure (35%) - Already screened (8.8%)		
Sawadogo et al <sup>28</sup>	840	- Knowledge of CC (64.2%) - Knowledge of HPV (8,5%) - CC Prevention (30.95%) - Contamination mode (19.9%) - Already screened (11.07%)		
Diendéré et al <sup>32</sup>	2293	- Already screened (6.2%)		

**Legend:** STI: Sexually Transmitted Infections; CC: cervical cancer; KAP: knowledge, attitudes and practices

### Genotypic coverage of Human Papillomavirus vaccines in Burkina Faso

HPV vaccination remains one of the most effective methods of primary prevention of cervical cancer. The genotypic coverage rate of HPV vaccines is a factor that guides the choice of vaccine among the three licensed

vaccines available on the market. Based on a pooled analysis of the studies collected, Figure 7 shows the genotype coverage rate of the three vaccines. The nonavalent vaccine Gardasil 9 covers 47.52% of the genotypes reported in this review. The overall number of genotypes not covered by available vaccines is 52.48%.



HPV vaccine name	HPV genotypes covered	Frequency of coverage in the general population n (%)
Bivalent (Cervarix)	16/18	461 (08.41%)
Quadrivalent (Gardasil)	6/11/16/18	573 (10.45%)
Nonavalent (Gardasil 9)	6/11/16/18/31/33/45/52/58	1468 (26.79%)
HR-HPV genotypes not covered by vaccines	35/39/41/49/51/53/56/59/66/68/82	1621 (29.58%)

**Figure 7.** Genotype coverage of Human Papillomavirus vaccines in the general population

### Discussion

In Burkina Faso, initiatives are underway to combat cervical cancer. However, up-to-date epidemiological data on HPV and cervical cancer are needed to assess the potential impact of these initiatives on cervical cancer prevention, and to identify effective strategies.

Based on 24 studies, this systematic review highlights the epidemiological profile of HPV and the prevention of cervical cancer among sexually active women in Burkina Faso.

This study shows poor coverage of epidemiological data on HPV and cervical cancer. Only studies from three of the 13 regions in Burkina Faso were included in this

review. The majority of studies were carried out in the “Centre” region, more precisely in the city of Ouagadougou. Despite this low coverage, we noted a high prevalence of HPV and a varied distribution, both genotypically and geographically in the regions, covered by the study. This prevalence varied from 20.6 to 97.59%, depending on the type of population. It was close to the results of other studies, which reported 8.9% and 81.8% in women from the general West African population<sup>3</sup> and 10.7 to 97.2% in women from sub-Saharan Africa<sup>16</sup>. It was high among women living with HIV, cervical cancer, sex workers and women infected with the hepatitis virus, with respective prevalences of 45.77%, 69.39%, 59.84% and 28%. Like all sexually transmitted infections (STIs), sex workers are exposed to

the HPV infection. With high unemployment, sex work remains a common practice in our countries, and it is not without risks, as it is a key factor in the high rate of STIs in sub-Saharan African countries. Studies on HPV prevention among sex workers in African countries reveal a much higher rate than ours<sup>42-45</sup>. This could be explained by the number of daily clients, the means of protection used by these sex workers, and also by the technique used for HPV genotyping in the various studies.

Immunodepression due to HIV infection is a major public health problem. However, according to UNAIDS, women and girls account for the majority of new HIV infections. Women living with HIV have a higher prevalence of genital high-risk oncogenic HPV (HR-HPV) infection than those in the general population<sup>46</sup>. They are more likely to be infected with several types of HR-HPV and have greater persistence of infection<sup>47</sup>. Nweke *et al* in 2013, Yakub *et al* in 2019 and Zinzendorf *et al* in 2022 reported a prevalence closer to our results<sup>48-50</sup> and Mkunde *et al* in 2023 recorded a prevalence of 50.9%, similar to our results<sup>51</sup>. In contrast, studies by Nyasenu *et al.* in 2019 recorded a prevalence of 22.2%<sup>52</sup>. This apparently reduced prevalence could be explained by the effectiveness of antiretroviral treatment and the competition among the viruses mentioned in some studies<sup>53,54</sup>.

Cervical cancer and CIN2/3 precancerous lesions are mostly caused by HR-HPV. The prevalence recorded in the present review was close to the 75.3% reported by Kelly *et al.* in 2017 in Burkina Faso<sup>46</sup>. Human Papillomavirus prevalence among women with unknown risk factors for HPV infection, based on collective analysis of data from the three regions of Burkina Faso, was 34.01%. These results corroborate those of Mkunde *et al* in 2023 who recorded 38.1%, 34% recorded by Seyoum *et al* in sub-Saharan Africa<sup>16</sup>, and the 33.6% by Ouedraogo *et al.* in West Africa<sup>3</sup>. For women in the general population of Burkina Faso, the pooled analysis shows an overall prevalence of 40.73%. This result is significantly higher than that of Ouédraogo *et al.* who recorded a prevalence of 28.6% in West Africa, and closer to that of Ogembo *et al.* who reported a prevalence of 50.5% in 2015 in a meta-analysis in Africa<sup>55</sup>. This variation in the prevalence of HPV infection could be explained by the difference in sample sizes and also the inclusion of cases of precancerous lesions, cervical cancer and female populations at high risk of HPV infection in some studies.

In this review, the HPV genotypes included were HR-HPV (16/18/31/33/35/39/45/51/52/56/58/59/66/68/82) and LR-HPV (6/11/43/44,55/54/69,71/70). It showed that the prevalence and distribution of these HPV genotypes in Burkina Faso varied according to region and study population, and 2232 of the 5480 women were infected with at least one of these HPV genotypes. The most frequent high-risk oncogenic genotypes were HPV18/52/56/35/31/58/66/59/51/16/39/45/68/33/53/82/41 in descending order for the general population. This distribution of HR-HPV genotypes could confirm the observations of Ouedraogo *et al.* in 2023, who reported that the distribution of HR-HPV in West Africa varied from country to country. Seyoum *et al.* in

2022 reported that HR-HPV genotypes in West African countries were distributed differently from those in Southern and East African countries. Ouedraogo *et al.* in 2023 explained this difference in distribution by human genetic factors. The second most frequent genotype in this review, HPV52, also attracted the attention of Ouedraogo *et al.* in 2023, who reported that it was common among the five main genotypes in most of the West African countries included in their study. Unfortunately, it is not included in the licensed quadrivalent cervical cancer vaccine used in Burkina Faso, despite the fact that it has been included in the cases of cervical cancer reported in some African studies<sup>56-58</sup>. Thanks to the support of Technical and Financial partners, the quadrivalent vaccine Gardasil 4 has been integrated into Burkina Faso's Expanded Program on Immunization. This commitment would require particular attention to the coverage rate of the most frequent genotypes in Burkina Faso. Our pooled analysis of studies carried out on HPV-infected women in Burkina Faso revealed that the nine most frequent genotypes were HPV18/52/56/35/31/58/66/59/51. Only HPV18 of the 9 is covered by Gardasil 4, and four genotypes (HPV18/52/31/58) are covered by Gardasil 9.

In this study, the bivalent Cervarix and quadrivalent Gardasil 4 vaccines were poorly covered in infected women. However, 14.92% of genotypes would be protected by Cervarix, 18.55% by quadrivalent Gardasil 4 and 52.48% not covered by the three vaccines. Only Gardasil 9 showed greater genotypic coverage. This low genotypic coverage had been reported in studies in Burkina Faso<sup>1,19,27</sup> and West Africa<sup>3,59</sup> despite the efficacy of these vaccines on the genotypes covered<sup>60</sup>. Despite these vaccines having some cross-protection against other less common HR-HPV types, as reported by some authors<sup>61</sup>, our results advocate the introduction of Gardasil 9 vaccine in Burkina as an excellent way to prevent cervical cancer, a silent killer of the female population.

In addition to the primary prevention provided by vaccination in the fight against cervical cancer, other approaches such as secondary prevention and risk factor prevention are key to cervical cancer prevention<sup>62,63</sup>. Most of the time, effective screening programs will detect precancerous lesions<sup>64</sup>. Despite the cost of cervical smear screening, WHO has authorized the detection of lesions by VIA/VILI in developing countries<sup>65</sup>. According to WHO's strategy to eliminate cervical cancer by 2030, screening remains the key factor. In Burkina Faso, VIA/VILI screening is free of charge in health centers. Despite this, WHO estimates that in low-resource countries, only 5% of women are screened for cervical cancer, compared with 40-50% in developed countries<sup>66</sup>. The incidence and mortality of CC in Burkina Faso is still close to the West African average, which remains among the highest in the world<sup>11,67</sup>. Women's level of knowledge, practices and attitudes compromise the Country's efforts in this fight. Indeed, analysis of studies on knowledge, attitudes and practices in this review reveals a low level of knowledge. Less than 60% of women in the various studies are aware of Cervical cancer. This lack of knowledge makes women vulnerable. This finding is in line with previous studies, which have also revealed low

awareness of cervical cancer in African countries<sup>28,64,68-78</sup>. This lack of awareness is at the root of low screening rates in African countries<sup>79,79-81</sup>. According to Chigbu et al. in 2017 in Nigeria, the use of community health educators for door-to-door education on cervical cancer prevention was associated with a significant increase in the uptake of cervical cancer screening and HPV vaccination<sup>82</sup>. According to WHO, community mobilization, health education and counseling are essential elements of an effective cervical cancer control program, as they help to ensure high vaccination coverage, high screening coverage and good treatment compliance. Health education ensures that women, their families and the community at large understand that cervical cancer can be prevented<sup>65</sup>.

Given the low level of knowledge about cervical cancer, prevention methods, risk factors and the low screening rate in Burkina Faso, health education about cervical cancer is still needed in sub-Saharan African countries, particularly Burkina Faso. The mapping of oncogenic HPV genotypes and the development of a vaccine covering these genotypes could be a breakthrough in the fight against cervical cancer in Africa.

### Study limits

The main objective of this review was to assess the epidemiological profile of HPV and the prevention of cervical cancer in Burkina Faso. It provides important insights into this infection in the female population, but the applicability of its findings is limited. Thus, this study presents limitations such as the diversity of HPV genotyping kits used in the different studies, the number of regions that have been the subject of HPV epidemiological studies with available data (three regions out of the thirteen regions of the country), the non-inclusion of low-risk genotypes in most of our studies included in this systematic review. Studies reporting indeterminate genotypes would also be a limitation for our study.

### Conclusion

In Burkina Faso, the fight against cervical cancer remains a major challenge, and HPV infection is a major health problem due to its high prevalence and involvement in the majority of cervical cancer cases. This prevalence varies from one region to another and depends on the type of

population. The distribution of oncogenic HPV genotypes responsible for the development of cervical cancer, varies according to the locality and population concerned in Burkina Faso. For more effective prevention of HPV infection and cancer, a large-scale vaccination program with Gardasil 9 would be beneficial. Large-scale national epidemiological studies are needed to determine the risk factors for HPV infections and progression to cervical cancer. In addition, it will be necessary to map HPV genotypes in cases of high-grade precancerous lesions and histologically confirmed cervical cancer, in order to make appropriate decisions for cervical cancer control. Women had insufficient knowledge of risk factors, prevention and screening for cervical cancer, which influenced their attitudes towards adherence to screening. A cervical cancer awareness program should focus on risk factors, prevention methods and the organization of screening campaigns. This information can be used to improve the planning and design of prevention interventions.

### Authors' contributions

Conceptualization: PZ and TS. Methodology: PZ, TS, RAO and JS participated in data collection. PZ, TS and RAO reviewed relevant articles; performed cluster analysis; participated in preparation of original project. Writing: PZ.

Revision and editing: PZ, TS, RAO, JS. All authors contributed to data interpretation and discussion and approved the final manuscript.

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### Conflict of interest

The authors declare that they have no conflicts of interest.

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## References

1. Maria H, Dana H, Françoise M, Michael P, Jürgen W. Human papillomaviruses in Western Africa: prevalences and risk factors in Burkina Faso. *Arch Gynecol Obstet.* 2018;298(4):789-796. doi:10.1007/s00404-018-4860-z
2. Bernard HU, Burk RD, Chen Z, van Doorslaer K, zur Hausen H, de Villiers EM. Classification of Papillomaviruses (PVs) Based on 189 PV Types and Proposal of Taxonomic Amendments. *Virology.* 2010;401(1):70-79. doi:10.1016/j.virol.2010.02.002
3. Ouedraogo RA, Kande A, Nadembega WMC, Ouermi D, Zohoncon TM, Djigma FW, Ouedraogo CMRN, Lompo OM, Simpore J. Distribution of high- and low-risk human papillomavirus genotypes and their prophylactic vaccination coverage among West African women: systematic review. *J Egypt Natl Cancer Inst.* 2023;35(1):39. doi:10.1186/s43046-023-00196-x
4. Duport N. Données épidémiologiques sur le cancer du col de l'utérus. *État Connaiss - Actual.* Published online January 1, 2008:34.
5. Ishida W, Singto Y, Yuenyao P, Tassaneeyakul W, Kanjanavirojikul N, Ishida T. Contribution of epigenetic risk factors but not p53 codon 72 polymorphism to the development of cervical cancer in Northeastern Thailand. *Cancer Lett.* 2004;210:205-211. doi:10.1016/j.canlet.2004.03.039
6. Muñoz N, Castellsagué X, Berrington de González A, Gissmann L. Chapter 1: HPV in the etiology of human cancer. *Vaccine.* 2006;24 Suppl 3:S3/1-10. doi:10.1016/j.vaccine.2006.05.115
7. Jain M, Yadav D, Jarouliya U, Chavda V, Yadav AK, Chaurasia B, Song M. Epidemiology, Molecular Pathogenesis, Immuno-Pathogenesis, Immune Escape Mechanisms and Vaccine Evaluation for HPV-Associated Carcinogenesis. *Pathogens.* 2023;12(12). doi:10.3390/pathogens12121380
8. Zohoncon TM, Ouedraogo TC, Brun LVC, Obiri-Yeboah D, Djigma WF, Kabibou S, Ouattara S, Gomina M, Yonli AT, Bazie VJTE, Ouedraogo C, Lompo O, Akpona SA, Simpore J. Molecular Epidemiology of High-Risk Human Papillomavirus in High-Grade Cervical Intraepithelial Neoplasia and in Cervical Cancer in Parakou, Republic of Benin. *Pak J Biol Sci PJBS.* 2016;19(2):49-56. doi:10.3923/pjbs.2016.49.56
9. Clifford GM, Gallus S, Herrero R, Muñoz N, Snijders PJF, Vaccarella S, Anh PTH, Ferreccio C, Hieu NT, Matos E, Molano M, Rajkumar R, Ronco G, de Sanjosé S, Shin HR, Sukvirach S, Thomas JO, Tunsakul S, Meijer CJLM, Franceschi S, IARC HPV Prevalence Surveys Study Group. Worldwide distribution of human papillomavirus types in cytologically normal women in the International Agency for Research on Cancer HPV prevalence surveys: a pooled analysis. *Lancet Lond Engl.* 2005;366(9490):991-998. doi:10.1016/S0140-6736(05)67069-9
10. Bhatla N, Aoki D, Sharma DN, Sankaranarayanan R. Cancer of the cervix uteri. *Int J Gynecol Obstet.* 2018;143:22-36. doi:10.1002/ijgo.12611
11. Ferlay J, Colombet M, Soerjomataram I, Mathers C, Parkin DM, Piñeros M, Znaor A, Bray F. Estimating the global cancer incidence and mortality in 2018: GLOBOCAN sources and methods. *Int J Cancer.* 2019;144(8):1941-1953. doi:10.1002/ijc.31937
12. Marie Tebeu P, Saint Saba Antaon J, Adjeba M, Pikop F, Tsuala Fouogue J, Ndom P. Connaissances, attitudes et pratiques des professionnels de santé sur le cancer du col de l'utérus au Cameroun: *Santé Publique.* 2021;Vol. 32(5):489-496. doi:10.3917/spub.205.0489
13. Mboumba Bouassa RS, Prazuck T, Lethu T, Meye JF, Bélec L. Cervical cancer in sub-Saharan Africa: an emerging and preventable disease associated with oncogenic human papillomavirus. *Med Sante Trop.* 2017;27(1):16-22. doi:10.1684/mst.2017.0648
14. Debrah O, Agyemang-Yeboah F, Donkoh ET, Asmah RH. Prevalence of vaccine and non-vaccine human papillomavirus types among women in Accra and Kumasi, Ghana: a cross-sectional study. *BMC Womens Health.* 2021;21(1):372. doi:10.1186/s12905-021-01511-1
15. Donkoh ET, Asmah RH, Agyemang-Yeboah F, Dabo EO, Wiredu EK. Prevalence and distribution of vaccine-preventable genital human papillomavirus (HPV) genotypes in Ghanaian women presenting for screening. *Cancer Control.* 2022;29:10732748221094721. doi:10.3389/fpubh.2022.890880
16. Seyoum A, Assefa N, Gure T, Seyoum B, Mulu A, Mihret A. Prevalence and Genotype Distribution of High-Risk Human Papillomavirus Infection Among Sub-Saharan African Women: A Systematic Review and Meta-Analysis. *Front Public Health.* 2022;10:890880. doi:10.3389/fpubh.2022.890880
17. Ogochukwu TN, Akabueze J, Ezeome IV, Aniebue UU, Oranu EO. Vaccination against human papilloma virus in adolescent girls: mother's knowledge, attitude, desire and practice in Nigeria. *J Infect Preve Med.* 2017;5(151):2.
18. Moher D, Liberati A, Tetzlaff J, Altman DG. Preferred reporting items for systematic reviews and meta-analyses: The PRISMA statement. *Int J Surg.* 2010;8(5):336-341. doi:10.1016/j.ijsu.2010.02.007
19. Madeleine KK, Djénéba O, Mahoukèdè ZT, Traore Fatié Porzé Wilfried, Gnoumou Ouamini Pulchérie De Prisca, Alice OR, Théophile YA, Prosper B, Paul O, Clarisse OTW, Edwige YT, Akouélé KKP, Dorcas OY, Ouedraogo Charlemagne Marie Ragnag-Néwendé, Jacques S. Molecular epidemiology of human papillomavirus in pregnant women in Burkina Faso. *Biomol Concepts.* 2022;13(1):334-340. doi:10.1515/bmc-2022-0026
20. Djigma FW, Zohoncon TM, Douamba Z, Sorgho PA, Obiri-Yeboah D, Ouattara AK, Sagna T, Traore L, Ghilat-Avoid-Belem NW, Sanogo K, Sempore J, Yonli AT, Pietra V, Bisseye C, Ouedraogo C, Simpore J. Molecular Genotyping of Human Papillomavirus among HIV-infected and HIV- uninfected Women in Ouagadougou, Burkina Faso. *Burkina Faso.* 2020;8.
21. Ouedraogo CMR, Djigma FW, Bisseye C, Sagna T, Zeba M, Ouermi D, Karou SD, Pietra V, Buelli F,



- Ghilat-Avoid-Belem NW, Sanogo K, Sempore J, Moret R, Pignatelli S, Nikiema JB, Sempore J. Épidémiologie et caractérisation des génotypes de papillomavirus humain dans une population de femmes à Ouagadougou. *J Gynécologie Obstétrique Biol Reprod.* 2011;40(7):633-638. doi:10.1016/j.jgyn.2011.05.012
22. Salambanga C, Zohoncon TM, Traoré IMA, Ouedraogo RA, Djigma WF, Ouédraog C, Sempore J. High prevalence of high-risk human papillomavirus (HPV) infection among sexually active women in Ouagadougou. *Médecine Santé Trop.* 2019;29(3):302-305. doi:10.1684/mst.2019.0920
  23. Djigma FW, Ouédraogo C, Karou DS, Sagna T, Bisseye C, Zeba M, Ouermi D, Gnoula C, Pietra V, Ghilat-Avoid-Belem NW, Sanogo K, Sempore J, Pignatelli S, Ferri AM, Nikiema JB, Sempore J. Prevalence and genotype characterization of Human Papillomaviruses among HIV-seropositive in Ouagadougou, Burkina Faso. *Acta Trop.* 2011;117(3):202-206. doi:10.1016/j.actatropica.2010.12.007
  24. Zohoncon TM, Bisseye C, Djigma FW, Yonli AT, Compaore TR, Sagna T, Ouermi D, Ouédraogo CMR, Pietra V, Nikiéma JB, Akpona SA, Sempore J. Prevalence of HPV High-Risk Genotypes in Three Cohorts of Women in Ouagadougou (Burkina Faso). *Mediterr J Hematol Infect Dis.* 2013;5(1):e2013059. doi:10.4084/MJHID.2013.059
  25. Ouedraogo RA, Zohoncon TM, Guigma SP, Angèle Traore IM. Oncogenic human papillomavirus infection and genotypes characterization among sexually active women in Tenkodogo at Burkina Faso, West Africa. *Papillomavirus Res.* 2018;6:22-26. doi:10.1016/j.pvr.2018.09.001
  26. Ouedraogo RA, Zohoncon TM, Guigma SP, Ouattara AK, Sempore J. Prédominance du papillomavirus humain 56 dans une sous-population de femmes sexuellement actives à Garango, Centre-Est, Burkina Faso: Predominance of Human Papillomavirus 56 in a subpopulation of sexually active women in Garango, Central-East, Burkina Faso. *J Appl Biosci.* 2020;150(1):15499-15509.
  27. Ouedraogo RA, Zohoncon TM, Traore IMA, Ouattara AK, Guigma SP, Djigma FW, Obiri-Yeboah D, Ouedraogo C, Sempore J. Genotypic distribution of human oncogenic papillomaviruses in sexually active women in Burkina Faso: Central, Central-Eastern and Hauts-Bassins regions. *Biomol Concepts.* 2020;11(1):125-136. doi:10.1515/bmc-2020-0011
  28. Sawadogo B, Gitta SN, Rutebemberwa E, Sawadogo M. Knowledge and beliefs on cervical cancer and practices on cervical cancer screening among women aged 20 to 50 years in Ouagadougou, Burkina Faso, 2012: a cross-sectional study. *Pan Afr Med J.* 2014;18(175). doi:10.11604/pamj.2014.18.175.3866
  29. Compaore S, Ouedraogo CMR, Koanda S, Haynatzki G, Chamberlain RM, Soliman AS. Barriers to Cervical Cancer Screening in Burkina Faso: Needs for Patient and Professional Education. *J Cancer Educ Off J Am Assoc Cancer Educ.* 2016;31(4):760-766. doi:10.1007/s13187-015-0898-9
  30. Kagoné TS, Paré PG, Dembélé A, Kania D, Zida S, Bonané/Thiéba B, Ouédraogo A, Testa J, Simporé J, Méda N. Cervical cancer in the Hauts-Bassins region of Burkina Faso: Results of a screening campaign by visual inspection with acetic acid (VIA screening in Burkina Faso). *Afr J Reprod Health.* 2022;26(6):97-103.
  31. Tassebedo S, Winter CH, Traore IT, Ouattara A, Sawadogo M, Meda N. Cervical pre-cancer screening by visual inspection of the cervix after application of acetic acid in rural Burkina Faso: evaluation of women's knowledge, screening practice habits, acceptability and prevalence of risk factors and lesions in Bousse health district. *Pan Afr Med J.* 2023;45:135. doi:10.11604/pamj.2023.45.135.36933
  32. Diendéré J, Kiemtoré S, Coulibaly A, Tougri G, Ily NI, Kouanda S. Low attendance in cervical cancer screening, geographical disparities and sociodemographic determinants of screening uptake among adult women in Burkina Faso : results from the first nationwide population-based survey. *Rev DÉpidémiologie Santé Publique.* 2023;71(4):101845. doi:10.1016/j.respe.2023.101845
  33. Traore IMA, Zohoncon TM, Dembele A, Djigma FW, Obiri-Yeboah D, Traore G, Bambara M, Ouedraogo C, Traore Y, Sempore J. Molecular Characterization of High-Risk Human Papillomavirus in Women in Bobo-Dioulasso, Burkina Faso. *BioMed Res Int.* 2016;2016:7092583. doi:10.1155/2016/7092583
  34. Ouattara S, Somé DA, Dembélé A, Sanfo S, Zohoncon T, Ouattara AK, Bambara M, Dao B, Simporé J. Molecular Epidemiology of High-Risk Human Papilloma Virus Infection in Sexually Active Women at Bobo-Dioulasso University Teaching Hospital. *Open J Obstet Gynecol.* 2019;09(08):1178-1188. doi:10.4236/ojog.2019.98114
  35. Tovo SF, Zohoncon TM, Dabiré AM, Ilboudo R, Tiemtoré RY, Obiri-Yeboah D, Yonli AT, Dovo EE, Ouédraogo RA, Ouattara AK, Sorgho PA, Ouermi D, Djigma FW, Ouédraogo C, Sangaré L, Sempore J. Molecular Epidemiology of Human Papillomaviruses, Neisseria gonorrhoeae, Chlamydia trachomatis and Mycoplasma genitalium among Female Sex Workers in Burkina Faso: Prevalence, Coinfections and Drug Resistance Genes. *Trop Med Infect Dis.* 2021;6(2):90. doi:10.3390/tropicalmed6020090
  36. Ilboudo R, Traoré EM, Zohoncon TM, Ouédraogo RA, Traore MA, Bado P, Ouédraogo C, Djigma FW, Obiri D, Sakandé J, Sempore J, Ouedraogo C. Prevalence and Characterization of High-Risk Human Papillomavirus Genotypes among a Group of Sex Workers in Ouagadougou, Burkina Faso. *Burkina Faso.* Published online 2020.
  37. Ouédraogo C, Zohoncon TM, Traoré EMA, Ouattara S, Bado P, Ouedraogo CT, Djigma FW, Ouermi D, Obiri-Yeboah D, Lompo O, Akpona SA, Sempore J. Distribution of High-Risk Human Papillomavirus Genotypes in Precancerous Cervical Lesions in Ouagadougou, Burkina Faso. *Open J Obstet Gynecol.* 2016;6(4):196-204. doi:10.4236/ojog.2016.64025



38. Zohoncon TM, Bado P, Ouermi D, Traoré EMA, Ouattara S, Djigma FW, Traore IMA, Yonli AT, Obiri-Yeboah D, Ouédraogo C, Akpona SA, Lompo O, Simpore J. Molecular characterization of high-risk human papillomavirus genotypes involved in invasive cervical cancer from formalin-fixed, paraffin-embedded tissues in Ouagadougou, Burkina Faso. *Int J Curr Res*. 2016;8(09):39314-39318.
39. Ouédraogo E, Zohoncon TM, Bayala B, Bado P, Da RPC, Ouedraogo RA, Traoré IMA, Kuassi-Kpede PA, Ouédraogo S, Dovo EE, Traoré L, Yonli AT, Djigma FW, Lompo OM, Simpore J. HPV/HBV or HPV/HCV Co-Infections in Women Treated for Chronic Hepatitis at Hôpital Saint Camille in Ouagadougou, Burkina Faso. *Am J Mol Biol*. 2024;14(01):1-12. doi:10.4236/ajmb.2024.141001
40. Chikandiwa A, Kelly H, Sawadogo B, Ngou J, Pisa PT, Gibson L, Didelot MN, Meda N, Weiss HA, Segondy M, Mayaud P, Delany-Moretlwe S, Group on behalf of the HS. Prevalence, incidence and correlates of low risk HPV infection and anogenital warts in a cohort of women living with HIV in Burkina Faso and South Africa. *PLOS ONE*. 2018;13(5):e0196018. doi:10.1371/journal.pone.0196018
41. Didelot-Rousseau MN, Nagot N, Costes-Martineau V, Vallès X, Ouedraogo A, Konate I, Weiss HA, Van de Perre P, Mayaud P, Segondy M, Yerelon Study Group. Human papillomavirus genotype distribution and cervical squamous intraepithelial lesions among high-risk women with and without HIV-1 infection in Burkina Faso. *Br J Cancer*. 2006;95(3):355-362. doi:10.1038/sj.bjc.6603252
42. Adams AR, Nortey PA, Dorte BA, Asmah RH, Wiredu EK. Cervical human papillomavirus prevalence, genotypes, and associated risk factors among female sex workers in Greater Accra, Ghana. *J Oncol*. 2019;2019.
43. Tounkara FK, Tégoué I, Guédou FA, Goma-Matsétsé E, Koné A, Béhanzin L, Traoré S, Aza-Gnandji M, Keita B, Guenoun J, Coutlée F, Alary M. Human papillomavirus genotype distribution and factors associated among female sex workers in West Africa. *Grce M, ed. PloS One*. 2020;15(11):e0242711-e0242711. doi:10.1371/journal.pone.0242711
44. Diop-Ndiaye H, Beiter K, Gheit T, Sow Ndoye A, Dramé A, McKay-Chopin S, Tommasino M, Bouh Boye CS, Sylla B, Kane CT. Human Papillomavirus infection in senegalese female sex workers. *Papillomavirus Res Amst Neth*. 2019;7:97-101. doi:10.1016/j.pvr.2019.02.003
45. Morhason-Bello IO, Baisley K, Pavon MA, Adewole IF, Bakare RA, de Sanjosé S, Francis SC, Watson-Jones D. Oral, genital and anal human papillomavirus infections among female sex workers in Ibadan, Nigeria. *PloS One*. 2022;17(3):e0265269. doi:10.1371/journal.pone.0265269
46. Kelly HA, Ngou J, Chikandiwa A, Sawadogo B. Associations of Papillomavirus humain (HPV) génotypes avec néoplasie cervicale de haut grade (CIN2 ) dans une cohorte de femmes vivant avec le VIH au Burkina Faso et Afrique du Sud. *PloS One*. 2017;12(3):e0174117. doi:10.1371/journal.pone.0174117
47. Clifford GM, Gonçalves MAG, Franceschi S, Group for the H and HS. Human papillomavirus types among women infected with HIV: a meta-analysis. *AIDS*. 2006;20(18):2337. doi:10.1097/01.aids.0000253361.63578.14
48. Nweke IG, Banjo AAF, Abdulkareem FB, Nwadike VU. Prevalence of human papilloma virus DNA in HIV positive women in Lagos University Teaching Hospital (LUTH) Lagos, Nigeria. Published online 2013.
49. Yakub MM, Fowotade A, Anaedobe CG, Manga MM, Bakare RA, Abimiku BA. Human papillomavirus correlates of high grade cervical dysplasia among HIV-Infected women at a major treatment centre in Nigeria: a cross-sectional study. *Pan Afr Med J*. 2019;33:125. doi:10.11604/pamj.2019.33.125.17589
50. Zinzendorf NY, Asher CM, Sandrine AT, Pierre B, Timotée O, Agathe DK, Joseph LS, Thérèse AKB. Prevalence and Human Papillomavirus Genotypes Distribution among HIV-Positive and -Negative Young Women Aged 20 to 37-Years-Old with Cervical Cancer in Abidjan, Côte d'Ivoire. In: ; 2022. <https://api.semanticscholar.org/CorpusID:247861170>
51. Chachage M, Parikh AP, Mahenge A, Bahemana E, Mnkai J, Mbuya W, Mcharo R, Maganga L, Mwamwaja J, Gervas R, Kibuuka H, Maswai J, Singoei V, Iroezindu M, Fasina A, Esber A, Dear N, Imbach M, Crowell TA, Hern J, Song X, Hoelscher M, Polyak CS, Ake JA, Geldmacher C, AFRICOS Study Group. High-risk human papillomavirus genotype distribution among women living with and at risk for HIV in Africa. *AIDS Lond*. 2023;37(4):625-635. doi:10.1097/QAD.0000000000003437
52. Nyasenu YT, Gbeasor-Komlanvi FA, Ehlan A, Issa SAR, Dossim S, Kolou M, Yambiyio BM, Prince-david M, Salou M, Ekouevi DK, Dagnra A. Prevalence and distribution of Human Papillomavirus (HPV) genotypes among HIV infected women in Lomé, Togo. *PLoS ONE*. 2019;14. <https://api.semanticscholar.org/CorpusID:71714850>
53. Heard I, Palefsky JM, Kazatchkine MD. The Impact of HIV Antiviral Therapy on Human Papillomavirus (Hpv) Infections and Hpv-Related Diseases. *Antivir Ther*. 2004;9(1):13-22. doi:10.1177/135965350400900117
54. Bratcher LF, Sahasrabuddhe VV. The impact of antiretroviral therapy on HPV and cervical intraepithelial neoplasia: current evidence and directions for future research. *Infect Agent Cancer*. 2010;5:8. doi:10.1186/1750-9378-5-8
55. Ogembo RK, Gona PN, Seymour AJ, Park HSM, Bain PA, Maranda L, Ogembo JG. Prevalence of human papillomavirus genotypes among African women with normal cervical cytology and neoplasia: a systematic review and meta-analysis. *PloS One*. 2015;10(4):e0122488.
56. Kabir AS, Bukar M, Nggada HA, Rann HB, Gidado A, Musa AB. Prevalence of human papillomavirus genotypes in cervical cancer in Maiduguri, Nigeria. *Pan Afr Med J*. 2019;33.

- <https://api.semanticscholar.org/CorpusID:201172653>
57. Attoh S, Asmah R, Wiredu EK, Gyasi R, Tettey Y. Human papilloma virus genotypes in Ghanaian women with cervical carcinoma. *East Afr Med J*. 2010;87(8).
  58. Nartey Y, Amo-Antwi K, Hill PC, Dassah ET, Asmah RH, Nyarko KM, Agambire R, Konney TO, Yarney J, Damale N. Human papillomavirus genotype distribution among women with and without cervical cancer: Implication for vaccination and screening in Ghana. *PLoS One*. 2023;18(1):e0280437.
  59. Obiri-Yeboah D, Akakpo PK, Mutocheluh M, Adjei-Danso E, Allornuvor G, Amoako-Sakyi D, Adu-Sarkodie Y, Mayaud P. Epidemiology of cervical human papillomavirus (HPV) infection and squamous intraepithelial lesions (SIL) among a cohort of HIV-infected and uninfected Ghanaian women. *BMC Cancer*. 2017;17(1):688. doi:10.1186/s12885-017-3682-x
  60. Louie KS, De Sanjose S, Mayaud P. Epidemiology and prevention of human papillomavirus and cervical cancer in sub-Saharan Africa: a comprehensive review. *Trop Med Int Health*. 2009;14(10):1287-1302. doi:10.1111/j.1365-3156.2009.02372.x
  61. Safaeian M, Kemp TJ, Pan DY, Porras C, Rodriguez AC, Schiffman M, Cortes B, Katki H, Wacholder S, Schiller JT, Gonzalez P, Penrose K, Lowy DR, Quint W, van Doorn LJ, Herrero R, Hildesheim A, Pinto LA. Cross-protective vaccine efficacy of the bivalent HPV vaccine against HPV31 is associated with humoral immune responses. *Hum Vaccines Immunother*. 2013;9(7):1399-1406. doi:10.4161/hv.24340
  62. Diallo MH. Connaissances, attitudes et pratiques des femmes en matière de dépistage du cancer du col de l'utérus (CCU) au centre médical communal de Coronthie Conakry- Guinée. *J SAGO Gynécologie – Obstétrique Santé Reprod*. 2021;22(1). Accessed February 13, 2024. <http://www.jsago.org/index.php/jsago/article/view/88>
  63. OMS. Vaccins contre les papillomavirus humains: note de synthèse de l'OMS, mai 2017. May 12, 2017:28.
  64. Sampson CN, Nkpeebo SD, Degley TA. Knowledge, attitude and health beliefs on cervical cancer screening in Ajumako-Eyan-Essiam District, Ghana. *Can Oncol Nurs J Rev Can Nurs Oncol*. 2021;31(3):285-290. doi:10.5737/23688076313285290
  65. OMS. *La lutte contre le cancer du col de l'utérus: guide des pratiques essentielles*; 2017. Accessed January 7, 2021. <http://apps.who.int/iris/bitstream/10665/254713/1/9789242548952-fre.pdf?ua=1>
  66. OMS. Stratégie mondiale en vue d'accélérer l'élimination du cancer du col de l'utérus en tant que problème de santé publique. Accessed July 6, 2024. <https://www.who.int/fr/publications/i/item/9789240014107>
  67. Bray F, Laversanne M, Sung H, Ferlay J, Siegel RL, Soerjomataram I, Jemal A. Global cancer statistics 2022: GLOBOCAN estimates of incidence and mortality worldwide for 36 cancers in 185 countries. *CA Cancer J Clin*. 2024;74(3):229-263. doi:10.3322/caac.21834
  68. Perrin A. Connaissance des femmes sur les modalités de prévention et de dépistage du cancer du col de l'utérus. Published online 1991.
  69. Ketevei AA, Logbo-Akey E, Andele A, Adou R, Tipoh AA, Aboubakari AS, Akpadza K. Connaissances, attitudes et pratiques des femmes en âge de procréer sur le cancer du col de l'utérus à l'hôpital de Bé (Togo): Knowledge, attitudes and practices of women of childbearing age concerning cervical cancer at be hospital (Togo). *J Rech Sci L'Université Lomé*. 2023;25(3):175-181.
  70. Bouslah S, Soltani MS, Ben Salah A, Sriha A. Connaissances, attitudes et pratiques des femmes tunisiennes en matière de dépistage du cancer du sein et de celui du col de l'utérus. *Psycho-Oncol*. 2014;8(2):123-132. doi:10.1007/s11839-014-0460-8
  71. Olubodun T, Odukoya OO, Balogun MR. Knowledge, attitude and practice of cervical cancer prevention, among women residing in an urban slum in Lagos, South West, Nigeria. *Pan Afr Med J*. 2019;32(130):130-130. doi:10.11604/pamj.2019.32.130.14432
  72. Ahmed SA, Sabitu K, Idris SH, Ahmed R. Knowledge, attitude and practice of cervical cancer screening among market women in Zaria, Nigeria. *Niger Med J J Niger Med Assoc*. 2013;54(5):316-319. doi:10.4103/0300-1652.122337
  73. Daka M, Ngoma C, Kalusopa VM, Banda Y, Chikwanda EK, Mulumba A. Knowledge, Attitudes and Perceptions Influencing Cervical Cancer Screening among Women in Kitwe District, Copperbelt Province, Zambia. *Open J Obstet Gynecol*. Published online 2022. <https://api.semanticscholar.org/CorpusID:250157220>
  74. Nyambe A, Kampen JK, Baboo SK, Van Hal G. Knowledge, attitudes and practices of cervical cancer prevention among Zambian women and men. *BMC Public Health*. 2019;19:1-15.
  75. O O, J TG. Knowledge, attitudes and practices of cervical cancer screening among rural women in KwaZulu-Natal, South Africa. *Pan Afr Med J*. 2022;42. doi:10.11604/pamj.2022.42.188.26172
  76. Shu NE, Abiola AHO, Akodu BA, Bassey BA, Misago N. Knowledge, attitudes and preventive practices for human Papilloma virus infection among female sex workers in Lagos metropolis. *Pan Afr Med J*. 2020;36:278. doi:10.11604/pamj.2020.36.278.17912
  77. De Groot AS, Tounkara K, Rochas M, Beseme S, Yekta S, Diallo FS, Tracy JK, Teguate I, Koita OA. Knowledge, attitudes, practices and willingness to vaccinate in preparation for the introduction of HPV vaccines in Bamako, Mali. Seale H, ed. *PLoS One*. 2017;12(2):e0171631-e0171631. doi:10.1371/journal.pone.0171631
  78. Osborne A, James PB, Bangura C, Williams SMT, Wadsworth R, Lebbie A. Knowledge, prevalence, and risk factors for self-reported sexually transmitted diseases among University Students in Sierra Leone. *Sierra Leone J Biomed Res*. 2023;14(2):81-91.
  79. Dickson KS, Boateng ENK, Acquah E, Ayebeng C, Addo IY. Screening for cervical cancer among

- women in five countries in sub-saharan Africa: analysis of the role played by distance to health facility and socio-demographic factors. *BMC Health Serv Res.* 2023;23(1):61-61. doi:10.1186/s12913-023-09055-w
80. Biaye B, Gassama O, Dieme MEF, Toure Y, Cissé M, Wade M, Diop D, Assane B, Guèye M, Diouf AA, Moreau JC. Screening for Cervical Cancer by Visual Inspection with Acetic Acid (VIA) in Nabil Choucair Health Center—Dakar (Senegal). *Open J Obstet Gynecol.* Published online 2019. <https://api.semanticscholar.org/CorpusID:86405742>
  81. Gnatou GYS, Onivogui Z, Adoli LK, Gbeasor-Komlanvi FA, Sadio AJ, Konu YR, Tchankoni MK, Zida-Compaore WIC, Agbonon A, Ekouevi DK. Screening for Precancerous Cervical Lesions in Women of Reproductive Age in the Kara Region of Togo in 2022. *Open J Epidemiol.* 2023;13(04):293-305.
  82. Chigbu CO, Onyebuchi AK, Onyeka TC, Odugu BU, Dim CC. The impact of community health educators on uptake of cervical and breast cancer prevention services in Nigeria. *Int J Gynecol Obstet.* 2017;137(3):319-324. doi:10.1002/ijgo.12150