

Review Article Open Access

# The Prevalence of Cervical Human Papillomavirus Infection and the Most At-risk Genotypes Among Iranian Healthy Women: A Systematic Review and Meta-analysis

Mina Malary, Mahmood Moosazadeh<sup>1</sup>, Zeinab Hamzehgardeshi<sup>2</sup>, Mahdi Afshari<sup>3</sup>, Iman Moghaddasifar, Amin Afsharimoghaddam<sup>4</sup>

Department of Reproductive Health and Midwifery, Student Research Committee, Mazandaran University of Medical Sciences, Sari, Iran, <sup>1</sup>Health Sciences Research Center, Faculty of Health, Mazandaran University of Medical Sciences, Sari, Iran, <sup>2</sup>Department of Reproductive Health and Midwifery, School of Nursing and Midwifery, Mazandaran University of Medical Sciences, Sari, Iran, <sup>3</sup>Department of Community Medicine, Zabol University of Medical Sciences, Zabol, Iran, <sup>4</sup>Department of Surgery, Shoole of Medicine, Zabol University of Medical Sciences, Zabol, Iran

#### Correspondence to:

Dr. Mahmood Moosazadeh, Health Sciences Research Center, Faculty of Health, Mazandaran University of Medical Sciences, Sari, Iran. E-mail: mmoosazadeh1351@gmail.com

How to cite this article: Malary M, Moosazadeh M, Hamzehgardeshi Z, Afshari M, Moghaddasifar I, Afsharimoghaddam A. The prevalence of cervical human papillomavirus infection and the most at-risk genotypes among Iranian healthy women: A systematic review and meta-analysis. Int J Prev Med 2016;7:70.

#### **ABSTRACT**

**Background:** One of the most common sexual-transmitted infections among women is human papillomavirus (HPV) infection which is associated with genital cancers. Different studies in Iran reported various prevalences, and combining their results could be important for health policy makers. This study aims to determine the total prevalence of HPV infection as well as its related genotypes, particularly HPV16 and HPV18 among Iranian healthy women.

**Methods:** Searching the Scientific Information Database, Iranmedex, Magiran, Irandoc, PubMed, Google Scholar, Scopus, and ScienceDirect databanks using relevant keywords and excluding duplicates and irrelevant evidence followed by applying exclusion criteria and quality assessment, eligible articles were selected. Standard error of the prevalence was calculated based on binomial distribution. Random effects model was used because of the high heterogeneity among the results.

**Results:** Of 14 studies entered into the systematic review, 24 pieces of evidence reported the HPV prevalence among 7655 healthy and noncancerous women in different Provinces of Iran. Total prevalence of HPV, 9.4% (95% confidence interval [CI]: 6.8–12.02); HPV16, 2.03% (95% CI: 1.3–2.8); HPV18, 1.7% (95% CI: 0.9–2.5); and other genotypes of HPV, 5.3% (95% CI: 3.6–6.9) were estimated.

**Conclusions:** Our meta-analysis showed that the total prevalence of HPV and its high-risk genotypes (16 and 18) among healthy noncancerous Iranian women was very high.

Keywords: Cervix, genotype, human papillomavirus, Iran, polymerase chain reaction, prevalence

# Access this article online Quick Response Code: Website: www.ijpvmjournal.net/www.ijpm.ir DOI: 10.4103/2008-7802.181756

## INTRODUCTION

One of the most common sexually transmitted infections worldwide with a high pathogenicity occurs by human

This is an open access article distributed under the terms of the Creative Commons Attribution-NonCommercial-ShareAlike 3.0 License, which allows others to remix, tweak, and build upon the work non-commercially, as long as the author is credited and the new creations are licensed under the identical terms.

For reprints contact: reprints@medknow.com

papillomavirus (HPV)<sup>[1-4]</sup> which is considered as the main risk factor for cervical cancer. [4] Although most HPV infections are self-limited, some other nononcogenic types (especially 6 and 11) develop genital warts. Moreover, persistent infections (HPV oncogenic types) can lead to cervical, vaginal, penile, anal, oropharyngeal and probably skin and lung cancers.<sup>[5]</sup> HPV infection can develop as latent, subclinical, and clinical forms.<sup>[2]</sup> It also has a high tendency to squamous epithelial cells with clinical manifestations differ from benign hyperplasic proliferative epithelial lesions (wart, swelling, or dermal masses) to invasive cancers. [6] It takes 10 years from occurring HPV infection to developing malignancy.<sup>[7]</sup> HPV prevalence rate decreases with age, but relative contribution of persistent infections increases over time.[8] Therefore, there is a high burden of morbidity and mortality in the developing countries mainly due to the lack of organized screening programs. [9]

Out of 100 HPV genotypes, forty genotypes are genital types<sup>[10]</sup> dividing into two types: Low risk and high risk.<sup>[2]</sup> Among these genotypes, 13 (HPV16, HPV18, HPV31, HPV33, HPV35, HPV39, HPV45, HPV51, HPV52, HPV56, HPV58, HPV59, and HPV66) are carcinogenic (high-risk types) while other types are low risk or with unknown risk,<sup>[8]</sup> and HPV16 and HPV18 are two common types.<sup>[11]</sup> HPV16, HPV18, and HPV16/18 are identifiable in 50–60%, 10–20%,<sup>[8]</sup> and 70% of cervical tumor specimens, respectively.<sup>[3,7,8,12,13]</sup>

About 10–20% of women without cervical dysfunction are infected with HPV. The corresponding figures for sub-Saharan Africa, Eastern Europe, and Latin America are 24%, 21%, and 16%, respectively. [11] HPV infection is more common in developing countries (42.4%) than among developed countries (22.6%). [14] Approximately 233.9 million women in reproductive age worldwide, 32.03 million Asian women, and 3.71 million Iranian women are HPV infected. [11] According to the WHO reports, out of four women, three women develop HPV infection at least once in their lifetime. Therefore, HPV is a common infection can affect each woman and is not related to any specific populations. [10]

Comprehensive review of national and international databases identifies different evidence regarding HPV infection among Iranian women indicating various prevalence estimates. It is necessary to combine these results using statistical methods. This study aims to estimate the total prevalence of HPV infection and its most risky genotypes among Iranian women using meta-analysis.

#### **METHODS**

The current study is a systematic review and meta-analysis regarding estimation of HPV infection prevalence and its hazardous genotypes among Iranian women conducted by the review of literature.

#### Search strategy

To find published electronic studies until February 20, 2015, national (Scientific Information Database, Iranmedex, Magiran, and Irandoc) and international (PubMed, Google Scholar, Scopus, and ScienceDirect) databases were reviewed. The following keywords and their Farsi equivalents were applied during searching processes:

"Prevalence," "seroprevalence," "frequency," "seroepidemiology," "human papillomavirus," "HPV," "pap smear," "cervical cancer," "PCR," "polymerase chain reaction," "HPV16," "HPV18," "high-risk genotypes," "Iran."

Our search was performed between February 21 and 29, 2015. To increase the search sensitivity, we investigated the lists of references belonged to the reviewed studies. One of our research team members randomly evaluated this search and reported no omitting of any relevant study. We also interviewed some experts and research centers in the study field to find probable unpublished studies.

### **Study selection**

Full text or abstract of all articles, documents, and reports identified during our systematic search was extracted. We first exclude all duplicates. Then, irrelevant papers were removed after reviewing titles, abstracts, and full texts. To prevent reprint bias, investigating the results with regard to identification of any repeated study was taken into consideration.

## **Quality** assessment

After selection of papers based on the titles and contents, their quality assessment was carried out using special checklists<sup>[15]</sup> derived from STROBE checklist,<sup>[16]</sup> containing 12 questions regarding different aspects of methodology such as appropriate sample size and sampling strategy, study population, data collection methods and tools, variable definition and methods of deal with samples, statistical tests, study objectives, and illustration and reporting of the results according to the study aims. One score was assigned to each question and studies achieved at least eight scores<sup>[15]</sup> were eligible to enter into the final meta-analysis.

#### **Inclusion** criteria

All Persian and English studies passed the evaluation processes and obtained the required quality scores and also estimated the HPV infection prevalence as well as its high-risk genotypes among healthy and noncancerous Iranian women based on cervical samples were selected.

#### **Exclusion criteria**

Studies did not report the prevalence of HPV infection, its high risk genotypes or sample size, those carried out among men, cancerous women, abnormal pap smears or saliva samples, duplicates or repeated results, studies reported in congresses or meetings without full-text presentation, case—control or clinical trials whose results did not reliably estimate the prevalence rate, and finally, articles without minimum quality assessment

scores were excluded from the systematic review and meta-analysis.

#### **Data extraction**

Study title, first author name, study date, sample size and sampling design, total prevalence of HPV and its genotypes, place of study, language of publication, and mean age of studied women were extracted from each study and entered into Excel Spreadsheet.

# Meta-analysis

All data analysis was performed by STATA SE Version 11 software (Stata Corporation, College Station, TX, USA). Standard error of the prevalence was calculated according to the binomial distribution equation. Cochrane test and I<sup>2</sup> index were used to detect the degree of heterogeneity between the results of different studies. If Q statistics showed statistical significance (P < 0.1) or when  $I^2$  index was near to one, we considered that the heterogeneity between the results is considerable so that pooling the results by random effects model would be required. Accordingly, prevalence of HPV infection as well as its genotypes was estimated using random effects model. In addition, using Bayesian analysis, we adjusted all point estimates to control the random variations. Moreover, we conducted sensitivity analysis to determine studies with the most influence on the heterogeneity. At least 20% change in the  $I^2$  index following the exclusion of each study indicated a significant influence of that study results in the heterogeneity. We also illustrated point and pooled estimates as well as their 95% confidence intervals (CIs) using forest plots. In these graphs, box size referred to study weight and crossed line represented 95% CI.

#### **RESULTS**

Searching national and international databases, 4737 primary studies were identified. Of them, 286 articles were duplicated; 4065 irrelevant articles were excluded after advanced search; 256 papers were omitted following title and abstract review; 107 irrelevant studies were removed after full-text review, and finally, 11 articles did not satisfy the inclusion/exclusion criteria or did not achieve enough quality scores. During the review of references, two eligible studies were added to the systematic review process. Finally, 14 papers were entered into the meta-analysis [Figure 1 and Table 1].

Of 14 finally selected studies, 24 pieces of evidence were reported with regard to HPV infection prevalence among healthy (noncancerous) women's pap smear specimens from different Provinces in Iran (Mobinikeshe in 2013 in a Persian-written study reported HPV infection rate among women in 11 Provinces in Iran). These studies have been published from 2002 to 2014, 12 of which written in English. Type of the study, sampling method, age groups, and mean age of the participants were reported in three,

http://www.ijpvmjournal.net/content/7/1/70

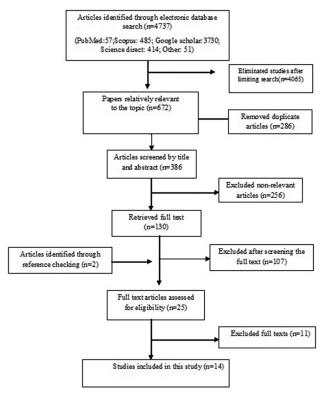


Figure 1: Literature search and review flowchart for selection of primary studies

six, seven, and four papers, respectively. Sample size had not been reported in one study and revealed after contact with corresponding author. All of the above studies used polymerase chain reaction as diagnostic method.

Total prevalences of HPV infection among women were differed from 0.63% in Eghbali study carried out in 2012 among 799 women to 36.2% in Eghbali study carried out among 47 women in 2013. Having adjusted by Bayesian analysis, the corresponding figures were 0.6 (Eghbali study) to 29.8% (Yousefzadeh study). Moreover, twenty studies reported the prevalence of HPV16 genotype among healthy pap smear specimens varied from zero in the study conducted by Mobinikeshe in 2013 among 232 samples to 29.8% in Eghbali study. Bayesian analysis limited these results to 0.1% (Eghbali) and 5.9% (Yousefzadeh). Prevalence of HPV-18 genotype was estimated in nine pieces of evidence from zero in the study carried out by Safaei in 2010 among 402 participants to 9.4% reported by Shahramian in 2011 among 265 women. These estimates changed after Bayesian analysis from 0.1% in Eghbali study to 3.4% in Shahramian study.

Among the entered studies, 19 pieces of evidence reported various HPV genotypes other than 16 and 18. Because of these high variations, reporting results in accordance with these genotypes was worthless. Prevalences of such genotypes varied between 0.4% in Eghbali study conducted among 799 women in 2012 and 21% in study carried out by Yousefzadeh among 851 participants in

Table 1: Baseline characteristics of included primary studies in meta-analysis of prevalence of human papillomavirus among Iranian healthy women

ID	First author	Publication year	Publication language	Sample size	Prevalence of HPV (%)			
					Total	Genotypes 16	Genotypes 18	Genotypes other
1	Allameh <sup>[17]</sup>	2012	Persian	180	25.5	3.9	3.3	12.8
2	Eghbali <sup>[18]</sup>	2012	English	799	0.63	0.1	0.1	0.4
3	Eghbali <sup>[19]</sup>	2013	English	47	36.2	29.8	6.4	-
4	Hamkar <sup>[20]</sup>	2002	English	44	9.1	-	-	9.1
5	Khodakarami <sup>[21]</sup>	2012	English	791	6.6	1.8	0.2	4.5
6	Afshar <sup>[22]</sup>	2013	English	350	24.6	-	-	-
7	Kesheh <sup>[23]</sup>	2013	Persian	377	4.8	1.6	-	3.2
8	Kesheh <sup>[23]</sup>	2013	Persian	308	4.2	1.6	-	2.6
9	Kesheh <sup>[23]</sup>	2013	Persian	257	5.4	1.6	-	3.8
10	Kesheh <sup>[23]</sup>	2013	Persian	171	2.3	1.7	-	0.5
11	Kesheh <sup>[23]</sup>	2013	Persian	226	3.1	1.3	-	1.8
12	Kesheh <sup>[23]</sup>	2013	Persian	231	3	0.9	-	2.1
13	Kesheh <sup>[23]</sup>	2013	Persian	125	4	0.8	-	3.2
14	Kesheh <sup>[23]</sup>	2013	Persian	274	2.2	0.4	-	1.8
15	Kesheh <sup>[23]</sup>	2013	Persian	205	2.4	0.5	-	1.9
16	Kesheh <sup>[23]</sup>	2013	Persian	232	9.5	0.4	-	9.1
17	Kesheh <sup>[23]</sup>	2013	Persian	232	3	0	-	3
18	Mohammadi <sup>[24]</sup>	2011	English	110	1.8	-	-	-
19	Moradi <sup>[25]</sup>	2011	English	378	20.1	5.8	4	10.3
20	Safaei <sup>[26]</sup>	2010	English	402	5.5	2	0	-
21	Shafaghi <sup>[27]</sup>	2014	English	851	31.1	7.3	2.8	21
22	Shahramian <sup>[28]</sup>	2011	English	265	33.4	9	9.4	15
23	Zandi <sup>[29]</sup>	2010	English	200	5.5	3.5	1.5	0.5
24	Jamali Zavarei <sup>[30]</sup>	2008	English	600	5.7	-	-	-

HPV=Human papillomavirus

2014. Bayesian analysis is adjusted these results as of 0.4% (Eghbali) and 18.8% (Yousefzadeh).

Due to the heterogeneity observed among the primary studies, we applied sensitivity analysis to identify studies mostly influenced the degree of heterogeneity. The study conducted by Eghbali among 799 women with HPV prevalence of 0.63 mostly affected the heterogeneity. Having excluded this study, the heterogeneity index changed from 97% to 96.3%. Continuing the sensitivity analysis showed that changes in the  $I^2$  index were inconsiderable. Therefore, meta-analysis was conducted using all studies entered into the final process.

As illustrated in Figures 2-5 and Table 2, prevalences of HPV infection and 16, 18, and other genotypes among Iranian healthy women using random effects model were estimated as of 9.4%, 2.03%, 1.7%, and 5.3%, respectively.

#### **DISCUSSION**

In this systematic review and meta-analysis, we estimated the total prevalence of HPV as well as its high-risk genotypes (HPV16 and HPV18) among Iranian healthy women as of 9.4%, 2.03%, and 1.7%, respectively.

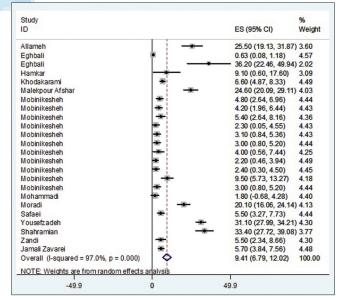


Figure 2: Prevalence of human papillomavirus overall in Iranian healthy women in per primary study and pooled estimate

In a study conducted in Romania among 1000 women aged between 17.3 and 57 years referred to Brasov Gynecology Hospital, prevalences of HPV infection with

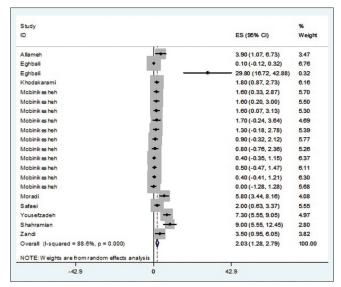


Figure 3: Prevalence of human papillomavirus genotype 16 in Iranian healthy women in per primary study and pooled estimate

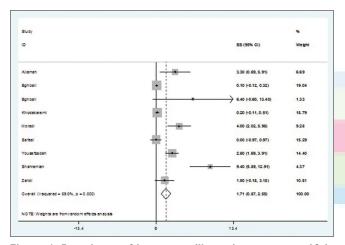


Figure 4: Prevalence of human papillomavirus genotype 18 in Iranian healthy women in per primary study and pooled estimate

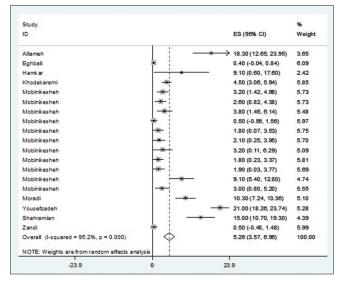


Figure 5: Prevalence of human papillomavirus other genotype in Iranian healthy women in per primary study and pooled estimate

35 different HPV genotypes were reported. HPV16 was the most common type (26.01%). Total prevalence of infection was significantly lower among women under 25. Moreover, HPV16 and HPV18 genotypes were more common among women with cervical abnormalities. Prevalences of high- and low-risk genotypes among these abnormal women were 48.2% and 3%, respectively.<sup>[51]</sup>

In a Turkish study carried out among 2161 normal and abnormal cervical cytological specimens, HPV infection prevalence was estimated as 12.6% most of whom carried HPV6 (5%), HPV18 (2.9%), and HPV16 (2.4%) genotypes.[32] Total prevalence of HPV among 725 women in Benin, Western Africa, was reported as 33.2%. Moreover, high-risk genotypes consisted about 88% of the infections, i.e. frequencies of HPV16 and HPV18 were 17.6% and 14.7%, respectively. The odds ratio of abnormal cytology among HPV-positive women was 2.9 times greater than that of HPV-negative women.[33] Among 60,775 18–79-year-old Korean women, prevalence of HPV infection was reported as 34.2%. Among HPV positives, 87.7% had only one type of infection while 12.3% had different types of HPV infection. The most common genotypes were HPV16 (26%), HPV52 (25.5%), HPV58 (12.3%), HPV18 (7.1%), and HPV56 (4.9%).[34]

In a study conducted in Portugal, total HPV infection prevalence was reported as 19.4%; most of them were 18-24-year-old women. About 16.5% of normal cytological specimens were infected. Approximately 76.5% of women were infected with HPV infection particularly with HPV16 (3.8%). Moreover, a statistically significant relationship (P < 0.001) was observed between high-risk HPV infection and abnormal cytology. [35] Prevalence of HPV infection among 899 Pakistani married women aged between 15 and 59 years was reported as 2.8%. Prevalence of HPV16 among women with or without abnormal cytology was 0.5% and 9.1%, respectively. In addition, number of partners, age difference between couples, other characteristics of husbands such as extra-marriage contacts, and their regular absents from the home were related with HPV infection prevalence. [36] In another study in Spanish among 2362 women, 34 different genotypes were identified. The most common genotypes were HPV16 (19.18%), HPV53 (11.26%), HPV58 (7.66%), and HPV18 (4.02%). HPV16 and HPV18 were detected among 24.3% and 5.1% of the high-risk infections, respectively. Both HPV16 and HPV18 were responsible for 30% of the high-risk infections in Spanish clinical centers.[37]

In a study carried out in China with a sample size of 4987, total prevalence of HPV and prevalence of high-risk HPV genotype were 13.3% and 10.2%, respectively. HPV52 (3.1%), HPV16 (2.5%), HPV58 (2.1%), HPV68 (1%), and HPV81 (0.9%) were the most common genotypes. [38] Moreover, in a study in Thailand, 29 different types of HPV were observed among 1662 women with average

Table 2: Pooled estimated of human papillomavirus prevalence and its high-risk genotypes in Iranian healthy women, according to the result of meta-analysis

Prevalence by	Sample size	Pooled estimate (%), 95% CI	Heterogeneity test		Heterogeneity (I <sup>2</sup> %)
			$\chi^2$ (Q)	P	
Overall HPV	7655	9.4 (6.8-12.02)	774.1	< 0.001	97
Genotype 16	6551	2.03 (1.3-2.8)	167.1	< 0.001	88.6
Genotype 18	3913	1.7 (0.9-2.5)	72.6	< 0.001	89
Other genotype	6146	5.3 (3.6-6.9)	377.8	< 0.001	95.2

HPV=Human papillomavirus, CI=Confidence interval

age of 43.4 years. Total prevalence of HPV and its prevalence among general population was 8.7% and 7.8%, respectively. Among HPV-positive samples, 44.8% were high-risk types. HPV infection was more common among 20–39-year-old women and then gradually decreased. [39] Roteli-Martins *et al.* in 2011 reported the total HPV infection among 3204 healthy 15–25-year-old women in Brazil, Canada, and the USA as of 26.6%. The prevalence of oncogenic HPV was 21.7% (25% in Brazil, 16.5% in Canada, and 19.1% in the USA). The most common types were HPV16 (5.2%), HPV51 (3.3%), HPV52 (3.3%), HPV31 (2.9%), HPV66 (2.3%), and HPV39 (2%). HPV infection was strongly associated with sexual behavior. [40]

According to the results of the current study, prevalence of HPV infection among healthy and noncancerous Iranian women was considerably lower than that reported among Roman, Brazilian, Canadian, American, Spanish, African, Portuguese, and Korean women. It was also slightly lower than the prevalences reported in Turkey and China. HPV infection rate in Iran was also higher than that of reported in Pakistan and Thailand. These differences might be due to different mean age of the participants, sampling selection methods, and variation in the prevalence of sexual behaviors and number of sexual partners. It should be noted that low age during marriage, first sexual contact in lower ages (<15), multiple sexual partners, and tobacco smoking were mostly related to HPV infection.[14] In addition, factors such as characteristics of the sexual partners, concomitance of sexual contact, and alcohol and drug abuse were the main risk factors of HPV infection.[3] In a meta-analysis, two-third of women with cervical cancer had HPV16 (51%) or HPV18 (16.2%) genotypes.[41]

In another meta-analysis, 16.88% (95% CI: 15.53%–18.31%) of patients with bladder cancer were infected with HPV. In addition, HPV infection increased the risk of bladder cancer more than 2.8 folds.<sup>[42]</sup>

HPV infection is a risk factor for invasive cervical cancer, and HPV16 and HPV18 genotypes are considered as carcinogenic factors for cervical cancer. Moreover, 31, 33, 35, 39, 45, 51, 52, 56, 58, 59, and 66 genotypes were also reported as high-risk genotypes of HPV infection. Among all HPV genotypes, 16 and 18 genotypes are the

most common types of HPV associated with cervical cancer. [23,43]

The heterogeneity observed between the results of the primary studies is one of the limitations of this systematic review and meta-analysis; therefore, we had to apply the random effects model to overcome this problem. Another limitation of the current study is that the most of the primary studies did not report factors associated with HPV infection. Even mean age of the participants did not report in many studies; therefore, we could not investigate the HPV infection risk factors during this meta-analysis. Further studies detecting the prevalence of all genotypes are suggested to be better applied for this reason.

Our meta-analysis presented evidence regarding HPV infection and its high-risk genotypes (16 and 18) among healthy Iranian women for policymakers. It should be noted that cervical cancer preventing programs have been planned according to control of HPV infection.

#### CONCLUSIONS

This meta-analysis showed that the prevalence of HPV and its high-risk genotypes among healthy Iranian women are considerable. Although it is less common in compare with the most other countries, the main part of cervical, breast, and bladder cancers among women can be attributed to this virus.

# Financial support and sponsorship Nil

#### **Conflicts of interest**

There are no conflicts of interest.

Received: 01 Aug 15 Accepted: 02 Apr 16 Published: 04 May 16

#### **REFERENCES**

- Jahdi F, Khademi K, Merghati Khoei E, Haghani H, Yazdanpanahi Z. Evaluation of the individual and medical factors associated with genital human papillomavirus in Iranian women. Scimetr 2014;2:e17495.
- Juckett G, Hartman-Adams H. Human papillomavirus: Clinical manifestations and prevention. Am Fam Physician 2010;82:1209-13.
- Javanbakht M, Gorbach PM, Amani B, Walker S, Cranston RD, Datta SD, et al. Concurrency, sex partner risk, and high-risk human papillomavirus

#### http://www.ijpvmjournal.net/content/7/1/70

#### International Journal of Preventive Medicine 2016, 7:70

- infection among African American, Asian, and Hispanic women. Sex Transm Dis 2010;37:68-74.
- Farzaneh F, Shirvani HE, Barouti E, Salehpour S, Khodakarami N, Alizadeh K. Knowledge and attitude of women regarding the human papillomavirus (HPV) infection, its relationship to cervical cancer and prevention methods. Med J Malaysia 2011;66:468-73.
- Reiter PL, Pendergraft WF 3<sup>rd</sup>, Brewer NT. Meta-analysis of human papillomavirus infection concordance. Cancer Epidemiol Biomarkers Prev 2010;19:2916-31.
- Shukla S, Bharti AC, Mahata S, Hussain S, Kumar R, Hedau S, et al. Infection of human papillomaviruses in cancers of different human organ sites. Indian J Med Res 2009;130:222-33.
- Azam F, Shams-ul-Islam M. Prevention of human papillomavirus infection with vaccines. J Pak Med Assoc 2010;60:676-81.
- Iftner T, Eberle S, Iftner A, Holz B, Banik N, Quint W, et al. Prevalence of low-risk and high-risk types of human papillomavirus and other risk factors for HPV infection in Germany within different age groups in women up to 30 years of age: An epidemiological observational study. J Med Virol 2010;82:1928-39.
- Garland SM, Smith JS. Human papillomavirus vaccines: Current status and future prospects. Drugs 2010;70:1079-98.
- Kawana K, Yasugi T, Taketani Y. Human papillomavirus vaccines: Current issues & future. Indian J Med Res 2009;130:341-7.
- Forman D, de Martel C, Lacey CJ, Soerjomataram I, Lortet-Tieulent J, Bruni L, et al. Global burden of human papillomavirus and related diseases. Vaccine 2012;30 Suppl 5:F12-23.
- Shikary T, Bernstein DI, Jin Y, Zimet GD, Rosenthal SL, Kahn JA. Epidemiology and risk factors for human papillomavirus infection in a diverse sample of low-income young women. J Clin Virol 2009;46:107-11.
- Krishna SK, Jethwa AS. Human Papillomavirus infections in adults and children.
   Am J Epidemiol Infect Dis 2013;1:11-9.
- Vinodhini K, Shanmughapriya S, Das BC, Natarajaseenivasan K. Prevalence and risk factors of HPV infection among women from various provinces of the world. Arch Gynecol Obstet 2012;285:771-7.
- Moosazadeh M, Nekoei-Moghadam M, Emrani Z, Amiresmaili M. Prevalence of unwanted pregnancy in Iran: A systematic review and meta-analysis. Int J Health Plann Manage 2014;29:e277-90.
- Elm EV, Altman DG, Egger M, Pocock SJ, Gotzsche PC, Vandenbroucke P, et al. The strengthening the reporting of observational studies in epidemiology (STROBE) statement: Guidelines for reporting observational studies. Prev Med 2007;45:247-51.
- 17. Allameh T, Moghim S, Farahbod F. Reviewing the prevalence of human papillomavirus (HPV) in married women aged 18-60 years with normal Pap smear referring to gynecology clinics in hospitals affiliated to Isfahan University of Medical Sciences, Iran. J Isfahan Med Sch 2012;29:2048-5.
- Eghbali SS, Amirinejad R, Obeidi N, Mosadeghzadeh S, Vahdat K, Azizi F, et al. Oncogenic human papillomavirus genital infection in Southern Iranian women: Population-based study versus clinic-based data. Virol J 2012;9:194.
- Eghbali M, Sadeghi F, Mirinargesi M. Frequency of human papillomavirus among pregnant women by PCR technique. Int J Mol Clin Microbiol 2013;2:285-8.
- Hamkar R, Mokhtari Azad T, Mohammadi M, Seyedirashti S, Severini A, Nategh R. Prevalence of human papillomavirus in Mazandaran province, Islamic Republic of Iran. East Mediterr Health J 2002;8:805-11.
- Khodakarami N, Clifford GM, Yavari P, Farzaneh F, Salehpour S, Broutet N, et al. Human papillomavirus infection in women with and without cervical cancer in Tehran, Iran. Int J Cancer 2012;131:E156-61.
- Afshar RM, Mollaie HR, Fazlalipour M, Arabzadeh SA. Prevalence and type distribution of human papillomavirus infection using the INNo-Lipa assay, Kerman, Southeast Iran. Asian Pac J Cancer Prev 2013;14:5287-91.
- Kesheh MM, Kaffashi A, Bagheri G, Karami MS, Mohammadi M, Naji A. Identification of human papillomavirus type 16 among Thinprep samples from 11 provinces of Iran. Iran J Obstet Gynecol Infertil 2013;16:22-8.
- Mohammadi A, Daneshbod K, Ghaffari N, Masoudian M, Rezvaninia S, Mohsenpour Z, et al. Evaluation of human papillomavirus infection among

- women using Pap smear and PCR in Shiraz, Iran. J Life Sci 2011;5:784-8.
- Moradi A, Bakhshandeh Nosrat S, Besharat S. Molecular epidemiology of high-risk types of human papillomaviruses (16, 18) in Pap-smear, the North East of Iran. Iran J Cancer Prev 2011;4:135-40.
- Safaei A, Khanlari M, Momtahen M, Monabati A, Robati M, Amooei S, et al. Prevalence of high-risk human papillomavirus types 16 and 18 in healthy women with cytologically negative Pap smear in Iran. Indian J Pathol Microbiol 2010;53:681-5.
- Shafaghi B, Jarollahi A, Yousefzadeh B, Ameri A, Moghadam S, Mostafavi M. Human papillomavirus prevalence and types among Iranian women attending regular gynecological visits. Rep Radiother Oncol 2013;1:73-9.
- Shahramian I, Heidari Z, Mahmoudzadeh-Sagheb H, Moradi A, Forghani F. Prevalence of HPV infection and high risk HPV genotypes (16, 18), among monogamous and polygamous women, in Zabol, Iran. Iran J Public Health 2011;40:113-21.
- Zandi K, Eghbali SS, Hamkar R, Ahmadi S, Ramedani E, Deilami I, et al. Prevalence of various human papillomavirus (HPV) genotypes among women who subjected to routine Pap smear test in Bushehr city (South West of Iran) 2008-2009. Virol J 2010;7:65.
- Jamali Zavarei M, Hamkar R, Dana VG, Delforoosh M, Shojamoradi MH, Gilani MM. Prevalence of HPV infection and its association with cytological abnormalities of Pap smears in Tehran. Iran J Public Health 2008;37:101-6.
- Moga MA, Irimie M, Oanta A, Pascu A, Burtea V. Type-specific prevalence of human papillomavirus by cervical cytology among women in Brasov, Romania. Asian Pac J Cancer Prev 2014;15:6887-92.
- Abike F, Bingöl B, Yılmaz A, Temizkan O, Tapısız ÖL, Dunder İ. HPV infection and HPV subtypes in normal and abnormal cervical cytology in Turkish women. J Virol Microbiol 2013;Article ID 640873:1-7.
- Piras F, Piga M, De Montis A, Zannou AR, Minerba L, Perra MT, et al. Prevalence of human papillomavirus infection in women in Benin, West Africa. Virol J 2011;8:514.
- Lee EH, Um TH, Chi HS, Hong YJ, Cha YJ. Prevalence and distribution of human papillomavirus infection in Korean women as determined by restriction fragment mass polymorphism assay. J Korean Med Sci 2012;27:1091-7.
- Pista A, de Oliveira CF, Cunha MJ, Paixao MT, Real O; CLEOPATRE Portugal Study Group. Prevalence of human papillomavirus infection in women in Portugal: The CLEOPATRE Portugal study. Int J Gynecol Cancer 2011;21:1150-8.
- Raza SA, Franceschi S, Pallardy S, Malik FR, Avan BI, Zafar A, et al. Human papillomavirus infection in women with and without cervical cancer in Karachi, Pakistan. Br J Cancer 2010;102:1657-60.
- Gomez-Roman JJ, Echevarria C, Salas S, González-Morán MA, Perez-Mies B, García-Higuera I, et al. A type-specific study of human papillomavirus prevalence in cervicovaginal samples in three different Spanish regions. APMIS 2009;117:22-7.
- Ye J, Cheng X, Chen X, Ye F, Lü W, Xie X. Prevalence and risk profile of cervical human papillomavirus infection in Zhejiang province, Southeast China: A population-based study. Virol J 2010;7:66.
- Chansaenroj J, Lurchachaiwong W, Termrungruanglert W, Tresukosol D, Niruthisard S, Trivijitsilp P, et al. Prevalence and genotypes of human papillomavirus among Thai women. Asian Pac J Cancer Prev 2010;11:117-22.
- Roteli-Martins CM, de Carvalho NS, Naud P, Teixeira J, Borba P, Derchain S, et al. Prevalence of human papillomavirus infection and associated risk factors in young women in Brazil, Canada, and the United States: A multicenter cross-sectional study. Int J Gynecol Pathol 2011;30:173-84.
- Clifford GM, Smith JS, Plummer M, Muñoz N, Franceschi S. Human papillomavirus types in invasive cervical cancer worldwide: A meta-analysis. Br J Cancer 2003;88:63-73.
- Li N, Yang L, Zhang Y, Zhao P, Zheng T, Dai M. Human papillomavirus infection and bladder cancer risk: A meta-analysis. J Infect Dis 2011;204:217-23.
- Serrano B, Alemany L, Tous S, Bruni L, Clifford GM, Weiss T, et al. Potential impact of a nine-valent vaccine in human papillomavirus related cervical disease. Infect Agent Cancer 2012;7:38.