HPV Vaccines: Current Status

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ABSTRACT

Cervical cancer is the fourth most common cancer in women worldwide, and the main cause is Human Papillomavirus (HPV) infection. The latest GLOBOCAN data reported an estimated 528,000 new cases and 266,000 deaths due to cervical cancer in 2012 (1). Besides, there were more than 600,000 new cases of HPV-related cancers, including anogenital and oral cancers in both genders. Current estimates indicate that every year 1686 women are diagnosed with cervical cancer and 663 die from the disease in Turkey (2). In 2013 a multicentric retrospective analysis held on 6,388 patients revealed that 25% of all women with normal and abnormal cytology had HPV infection (3).

In previous studies, deoxyribonucleic acid (DNA) of HPV was detected in the vast majority of patients with cervical dysplasia and carcinoma. Currently, it is considered to be prerequisite for the development of these cervical lesions, but needs a number of supplementary factors for the development invasive carcinoma. There are more than 100 types of HPV identified to be able to infect epithelial cells, but specific types have higher risk for oncogenic transformation. There are 15 high-risk anogenital types; HPV-16 is the most common type, and followed by HPV types 18, 45, 31, 33, 35, 52, and 58, whereas HPV types 51, 56, 39, 59, 68, 73, and 66 are much less common.

It is currently believed that there is a step-by-step development of cervical neoplasms; starting with a persistent HPV infection due to a high-risk type, followed by high-grade dysplasia, and finalized in invasive carcinoma. High-risk HPV types can also lead to the development of low-grade intraepithelial lesions (LSIL); however, these lesions substantially regress (4,5).

The discovery of the 15 HPV types which are the most common cause of cervical cancer has risen the idea of whether the development of cervical neoplasia can be prevented by HPV-based screening or primary or secondary prophylaxis with vaccines.

The quadrivalent and bivalent HPV vaccines were introduced in 2006 and 2007, respectively, for the female population. The bivalent vaccine included the most oncogenic HPV types, HPV-16 and HPV-18, which are responsible for 70% of all cervical cancers worldwide (6). The outer coat (L1) protein,
a virus-like particle, of these types were introduced into the vaccine in order to prevent cancers of the cervix, vulva, vagina, anus, penis, and oropharynx (7). In addition to HPV-16 and HPV-18, the quadrivalent vaccine also contains L1 proteins of HPV-6 and HPV-11, which are the leading cause of genital warts as well as recurrent respiratory papillomatosis (8). Studies have shown >90% efficacy for both vaccines for the prevention of cervical intraepithelial neoplasia (CIN), vulvar intraepithelial neoplasia (VIN), and vaginal intraepithelial neoplasia (VaIN) that were developed due to targeted HPV types, and for quadrivalent vaccine also for the prevention of genital warts (9). The most outrageous finding is that the protection of these vaccines has been observed to continue for at least 10 years based on antibody decay rates in modeling studies (10).

It should be kept in mind that these vaccines are only efficient for the aim of prophylaxis, and therefore not effective in the treatment of existing infections. For this reason, the target population of the vaccines are preadolescent girls that have never had a sexual intercourse. The safety profile in this population was evaluated after approximately 250 million doses of HPV vaccines were administered by the year 2015 worldwide, and no significant side effect was observed (11,12).

**Efficacy and Safety**

The efficacy of vaccines can be specified by their effectiveness on decreasing the rates of HPV infection and related diseases.

Markowitz et al. reported that HPV prevalence was reduced to 5.1% in females between 14 and 19 years from a rate of 11.5% with the administration of HPV vaccines (13). The vaccine coverage rate in this study was 32%. Similarly, The National Surveys of Sexual Attitudes and Lifestyles from the United Kingdom, in which the coverage rate was 61.5%, revealed the prevalence of HPV-16 and HPV-18 was reduced to 5.8% in women between 18–20 years from a rate of 11.3% (14). A number of other studies also showed the effectiveness of HPV vaccines on decreasing the rate of HPV infections and genital warts. A systemic review of these studies reported that HPV infections and anogenital warts were significantly reduced with the use of HPV vaccine, and also stated that the coverage of the vaccine was ≥50% (15).

An Australian study was the first to report the early results of the effect of a population-based HPV vaccine program on the development of premalignant lesions (16).

In this study, the rates of CIN 2-3 lesions were significantly reduced in females under 18 years of age with the introduction of HPV vaccines. There are four phase III trials in which the clinical efficacy of the HPV vaccine was studied; PATRICIA and Costa Rica trials for the bivalent, and FUTURE I and II trials for the quadrivalent vaccine (17-19). The efficacy of both vaccines was found 96-100% in terms of preventing HPV 16 and 18-related CIN, carcinoma in situ and cervical cancer. Antibody titers for both vaccines were measured significantly higher than those achieved with natural HPV infection. The duration of these high titers was 5 years in the studies of the quadrivalent vaccine and 8.4 years in the studies of the bivalent vaccine with seropositivity rates of 98.8% and 100%, respectively. This duration of preventing HPV infection and related disease may be longer when the final results of these studies are reported. However, current data reveals high rates of immunogenicity and adequate duration of protection without any need for a booster dose. The 7-year follow-up of the VIVIANE study revealed that HPV 16-18 vaccine is still successful in protecting women older than 25 years against infections, cytological abnormalities, and lesions associated with HPV 16-18 as well as HPV 31 and HPV 45 (20).

The WHO and CDC have confirmed the safety of HPV vaccines after >250 million doses were administered worldwide; however, they have also recommended a much longer duration for detecting whether there would be any adverse events. Although no adverse events were encountered in pregnant women that were inadvertently vaccinated, administration of HPV vaccines in pregnant women is not recommended (21).

**Alternative Strategies for Dosing**

The primary dose schedule was held as three doses; at months 0, 1 or 2 (for quadrivalent and bivalent, respectively), and 6, and both are currently administered in 3 doses for an optimized immune response. Both HPV vaccines contain virus-like particles which are highly immunogenic. No cut-off value for antibody levels was detected for protection against HPV infection. However, higher titers of antibodies were observed in adolescents than needed which are probably the main reason for higher rates of protection in adult women. With the aim of reducing the cost of vaccination, studies comparing 3-dose schedules to two-dose schedules (in months 0 and 6) were held for both bivalent and quadrivalent vaccines. A number of trials showed equivalent immunogenicity in two-dose schedules in young adolescents compared to three-dose schedules in adult women measured by antibody titers (22,23).

However, memory responses of T and B-cells were found lower with the two-dose schedule for the quadrivalent vaccine compared to three-dose schedule (24). Besides, although there are no data about the statistical significance, the antibody titers against HPV-18 and HPV-16 were lower after 24 months and 36 months, respectively, in adolescents that underwent two-dose schedule compared to the ones that received three doses of vaccination.

The Costa Rica trial was the first to show that two doses, or even one dose, of the bivalent HPV vaccine was successful in the protection against HPV infection (25,26). However, the long-term protection of the HPV vaccine was maintained in all women even they are not seropositive, probably with a similar reason to what is accepted for hepatitis B vaccine (27,28).

In 2014, based on these findings, World Health Organization (WHO) approved the two-dose schedule of HPV
vaccines in females younger than 15 years of age, with ≥6 month-intervals (29). Since the approval of the WHO Switzerland, Quebec and British Columbia of Canada, the United Kingdom, France, Spain, The Netherlands, South Africa, and Chile have adopted the two-dose schedule. However, while administering the two-dose schedules it is important to consider the relative cost-effectiveness of them, and risk management strategies should be kept under consideration for situations where they do not provide protection for an adequate time (30).

Vaccination of Males

The first identified HPV-related malignancy is cervical cancer; however, it has been clearly shown that HPV infection can also lead to anogenital, penile and oral cancers as well as anogenital warts in men. One of the main concerns about men is that by the disease load they carry, they can transmit the virus to women by sexual intercourse and thus, they are the indirect cause of the development of cervical cancer in women (31). Anal cancer has a higher risk of development particularly in men that have anal sexual intercourse with men (32). Development of genital warts and recurrent respiratory papillomatosis can also be observed in men. In order to prevent the fore-mentioned diseases, the concept of vaccination in men has arisen. But the main benefit from the vaccination of men would be for the sake of women, because herd protection significantly decreases the rate of the development of HPV infection in women.

Giuliano et al. included 4,065 healthy males between 16-26 years of age from 18 countries into a randomized, placebo-controlled, double-blinded trial in which they evaluated the efficacy of quadrivalent HPV vaccine on external genital lesions in men. No males had a history of anogenital warts, penile, perianal or perineal intraepithelial neoplasia or cancer. The efficacy of the vaccine was found 92.4% and 79% in 3,463 heterosexual males and in 602 males who had sex with males (MSM), respectively. When the MSM subgroup was evaluated for the primary end point of HPV-6, 11, 16 or 18-related anal intraepithelial neoplasia (AIN) grade 1 (including condyloma), 2 or 3, or anal cancer; the efficacy of the quadrivalent vaccine was found 77.5%, and a median 85% (range: 76.2%-100%) reduction was observed in the detection of HPV DNA at any time in the post-vaccine follow-up (33,34).

Male vaccination has been found cost-effective in a number of studies, particularly when the achieved coverage of female vaccination remains under 50%. Nevertheless, spending financial resources on female coverage is more rational in order to achieve an actual cost-effective status. (35)

Routine HPV vaccination for males has been recommended since 2011 in the United States; however, as it is not fully funded by the government and is mainly school-based, it did not achieve a coverage as high as in females (35% in males vs. 57% in females) (36).

Another country that recommends HPV vaccination in both genders is Australia. The coverage of male vaccination was no higher than 2% in 2006; however, it reached a rate of 54% in 2015 after a government funding was accepted for the program. There is an ongoing trial of HPV vaccination programs for both genders in Alberta and Prince Edward Island of Canada. Besides, an HPV vaccination program for men who have sex with men has been recently planned to be held in the United Kingdom.

Nine-valent HPV vaccine

As already mentioned, quadrivalent and bivalent vaccines have protection against specific HPV types that are responsible for the development of approximately 70% of all cervical cancers. Besides these, they also provide cross-protection for a number of other oncogenic types. However, studies on the efficacy and duration of cross-protection have not reported reliable data. With an aim of protection against a higher number of HPV types, the nine-valent vaccine which additionally includes L1 proteins of five other oncogenic HPV types, i.e. 31, 33, 45, 52, and 58, have been introduced with an expectation of protection against approximately 90% of all cervical cancers (Table 1).

Joura et al. randomized more than 14,000 females between 16 and 26 years to quadrivalent and nine-valent vaccines in a phase III trial in which the primary endpoints were rates of HPV infection and intraepithelial neoplasia (37).

Both vaccines were administered as three doses at months 0, 2 and 6. The overall efficacy of the nine-valent vaccine was found 96.7%, which was higher than the one achieved with the quadrivalent vaccine. The nine-valent vaccine also offered a satisfactory high rate of protection against CIN2 and CIN3, VIN2 and VIN3, and VaIN2 and VaIN3. Moreover, rate of systemic adverse events such as fever and nausea were similar with both vaccines. The only difference was the rate of pain at the injection site which was significantly higher in the nine-valent vaccine arm. It was also reported that antibody titers against HPV-6, 11, 16, and 18 did not decrease in the nine-valent vaccine compared to quadrivalent vaccine.

The Advisory Committee on Immunization Practices (ACIP) recommends the 9-valent vaccine and it has been approved by the FDA in 2014 for both genders between 11–12 years of age (38).

In individuals that were not vaccinated between this target age, catch-up vaccination is recommended until the age of 26 for both genders.

It is not currently recommended to administer an additional nine-valent HPV vaccine after a previous three-dose schedule of the quadrivalent or the bivalent HPV vaccine. However, it is considered safe to administer any kind of HPV vaccine in individuals of both genders that do not know the type of their previous HPV vaccine in an incomplete schedule they previously had.
**Conclusion**

The relation between HPV and development of cancer was and has been studied in various molecular and genetic trials as well as it has been shown in epidemiological data and clinical observations. In regard to the younger population it affects and years of life it causes to be lost, the main prophylaxis for cervical cancer is the prophylaxis of HPV. Further studies will enlighten the medical world about the efficacy of HPV vaccines in other HPV-related diseases in males and females. There is a tremendous hope for the development of new dose schedules and new vaccines which will be more efficacious in preventing various HPV-related cancers.

**References**


2. Human Papillomavirus and Related Cancers, Fact Sheet 2016. ICO Information Centre on HPV and Cancer (October 7, 2016)


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**Table 1: Characteristics of bivalent, quadrivalent, and nine-valent HPV vaccines (39)**

<table>
<thead>
<tr>
<th>HPV types included</th>
<th>Prevented Diseases</th>
<th>Efficacy</th>
</tr>
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<tbody>
<tr>
<td>Bivalent</td>
<td>HPV-16 and -18-related Cervical cancer CIN1-3 AIS</td>
<td>98.1 % for diseases related to HPV 16 and -18</td>
</tr>
<tr>
<td>Quadrivalent</td>
<td>HPV-6, -11, -16, and -18-related Cervical cancer Vulvar cancer Vaginal cancer, CIN1-3 AIS VIN 2-3 VaIN 2-3 PIN 1-3 Penile cancer Anogenital warts AIN Anal cancer</td>
<td>6, 11, 16, and 18</td>
</tr>
<tr>
<td>9-valent</td>
<td>HPV-6, -11, -16, -18, -31, -33, -45, -52, and -58-related Cervical cancer Vulvar cancer Vaginal cancer CIN 1-3 AIS VIN 2-3 VaIN 2-3 PIN 1-3 Penile cancer Anogenital warts AIN Anal cancer</td>
<td>&gt;99% for diseases related to HPV-6,-11,-16,-18,-31,-33,-45,-52,-58</td>
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**Table Notes:**

- **HPV:** Human Papilloma Virus, **CIN:** Cervical intraepithelial neoplasia, **AIS:** Adenocarcinoma in situ, **VIN:** Vulvar intraepithelial neoplasia, **VaIN:** Vaginal intraepithelial neoplasia, **PIN:** Penile intraepithelial neoplasia, **AIN:** Anal intraepithelial neoplasia.


34. Palefsky JM, Giuliano AR, Goldstone S, Moreira ED Jr,


39. ACOG-Committee on Adolescent Health Care and Immunisation Expert Work Group. Number 641-September 2015