Short Report

Effectiveness of the AS04-adjuvanted HPV-16/18 vaccine in reducing oropharyngeal HPV infections in young females—Results from a community-randomized trial

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We studied effectiveness of the AS04-adjuvanted HPV-16/18 (AS04-HPV-16/18) vaccine against human papillomavirus (HPV) oropharyngeal infections associated with the increase of head/neck cancers in western countries. All 38,631 resident adolescents from 1994 to 1995 birth cohorts of 33 Finnish communities were invited in this community-randomized trial (NCT00534638). During 2008–2009, 11,275 girls and 6,129 boys were enrolled in three arms of 11 communities each. In Arm A, 90% of vaccinated girls/boys, and in Arm B, 90% of vaccinated girls received AS04-HPV-16/18 vaccine. Other Arm A/B and all Arm C vaccinated participants received control vaccine. All Arm A participants and Arm B female participants were blinded to vaccine allocation. Oropharyngeal samples were analyzed from 4,871 18.5-year-old females who attended follow-up visit 3–6 years postvaccination. HPV DNA prevalence was determined by SPF-10 LiPA and Multiplex type-specific PCR. Total vaccine effectiveness (VE) was defined as relative reduction of oropharyngeal HPV prevalence in pooled Arms A/B HPV-vaccinated females vs. all Arm C females. VE against oropharyngeal HPV-16/18, HPV-31/45 and HPV-31/33/45 infections were 82.4% (95% confidence intervals [CI]: 47.3–94.1), 75.3% (95% CI: 12.7–93.0) and 69.9% (95% CI: 29.6–87.1), respectively. In conclusion, the AS04-HPV-16/18 vaccine showed effectiveness against vaccine and nonvaccine HPV-types oropharyngeal infections in adolescent females up to 6 years postvaccination.

Additional Supporting Information may be found in the online version of this article.
Key words: human papillomavirus, oral infection, oropharyngeal cancer, vaccine effectiveness

Abbreviations: Al: aluminum; AS04-HPV-16/18: AS04-adjuvanted HPV-16/18 vaccine; CI: confidence interval; GCP: good clinical practice; HBV: hepatitis B virus; HPV: human papillomavirus; hrHPV: high-risk human papillomaviruses; IARC: International Agency for Research on Cancer; MPL: monophosphoryl lipid A; n (%): number (percentage) of subjects reporting an event; N: number of subjects with available results; OPSCC: oropharyngeal squamous cell cancer; VE: vaccine effectiveness

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The incidence of oropharyngeal cancers due to human papillomavirus (HPV) infection is increasing in Western countries. However, the efficacy of HPV vaccines against oropharyngeal infection is not well documented. Based on a large community-randomized trial in Finland, the authors report high effectiveness (82.4%) of Cervarix vaccine against oropharyngeal HPV-16/18 infection and moderate effectiveness against non-vaccine types HPV-31/33/45 infection in female adolescents. The protective effects were observed up to 6 years after vaccination. These results provide further evidence that HPV vaccination holds the potential to reduce oral HPV infection, thereby offering protection against HPV-related head and neck cancers.

Introduction
Oncogenic, high-risk human papillomaviruses (hrHPV) cause worldwide 9% of cancers in females and 1% in males. More than 15 years ago, a retrospective case-series and a large nested case-control study showed that infection with HPV-16 is associated with a high risk of oropharyngeal squamous cell cancer (OPSCC), that develops 10 or more years after acquisition of the infection. Identification of HPV DNA in oropharyngeal and tonsillar tumor tissues have confirmed these observations. In 2009, the International Agency for Research on Cancer (IARC) recognized the causal role of HPV-16 in the etiology of cancers of the oropharynx and tonsil. The HPV association is most true for lymphoepithelial associated OPSCC-subsites infected with HPV.

In western countries, the incidence of HPV-positive OPSCC has increased very rapidly, virtually doubling in a decade since the 1970s. Such trends have not been observed in HPV-negative OPSCC. The former is clinically significantly more favorable and can be identified/monitored using HPV early antigen antibodies. Thus, HPV testing is important but prospects for screening of HPV-positive OPSCCs may not be optimal due to the lack of defined manageable precancerous lesions. Moreover, in vaccinated populations with low background prevalence the predictive values for any screening test will be demanding.

In the adolescent and young adult populations, the prevalence of oral HPV infections varies widely from 0.4% to 10% mostly depending on the method used for viral DNA detection. While HPV vaccination induces antibody levels at oral cavity that correlate with circulating antibody levels, the efficacy of prophylactic HPV vaccines against oral infections has not been extensively studied. Up to 88% reduction of oral hrHPV infections in vaccine recipients as compared to nonvaccinated adolescents was recently suggested but randomized trial data are scarce. Based on a large community-randomized trial we now report secondary objective results on the total vaccine effectiveness (VE) of the AS04-adjuvanted HPV-16/18 (AS04-HPV-16/18) vaccine against oropharyngeal HPV infections in adolescent females in Finland.

Materials and Methods
Community randomization
All the 33 randomized Finnish communities had more than 35,000 inhabitants. Each study arm comprised 4 high (>24%), 3 intermediate (24–20.5%) and 4 low (<20.5%) HPV-16/18 seroprevalence communities. HPV seroprevalence is a measure of the cumulative HPV incidence, and as such gives a valid estimate of the background HPV exposure. This reduced the coefficient of variation across arms, and increased statistical power.

Ethics
The primary HPV-040 trial (NCT00534638, clinicaltrial.gov) and its amendment for the collection of the oral gargle samples were approved by the ethical committee of the Pirkanmaa hospital district in 2007 and 2011. The trial was conducted in accordance with good clinical practice (GCP) and all applicable regulatory requirements including the Declaration of Helsinki. Participants aged <15 years provided written informed assent; written informed consent was obtained from the participants’ parents or legal representatives. Written informed consent was obtained from study participants aged 15 years in line with local regulations.

Study procedures
In the 2008–2009 school year, invitation letters were sent to the parents or legal guardians of all the 80,272 Finnish or Swedish speaking 1992–1995 born adolescents residents in the 33 communities identified by the Finnish Population Information System maintained by the Population Register Centre. The letters included trial information, parental and adolescents’ consent forms and a prepaid return envelope.

The vaccination (first three study visits) took place at the healthcare facilities of 250 municipal junior high schools of the study site communities. In Arm A, 90% of vaccinated participants received Cervarix (AS04-HPV-16/18; GSK) vaccine, and 10% received Engerix B vaccine (hepatitis B virus [HBV] vaccine; GSK). AS04 is an Adjuvant System containing monophosphoryl lipid A (50 μg MPL; produced by GSK) adsorbed on Aluminum salt (500 μg Al3+). In Arm B, 90% of vaccinated females received the AS04-HPV-16/18 vaccine, and 10% of vaccinated females and all vaccinated males received the HBV vaccine. The 90 and 10% proportions were receiver-blinded proportions to assess the herd effect. The blinding was maintained for girls and boys in Arm A and for girls in Arm B. In Arm C communities, all vaccinated participants received the HBV vaccine. Virtually all (99%) vaccinated participants received all three vaccine doses.

Both vaccinated and nonvaccinated females of the 1994–1995 birth cohorts from the study site communities were invited to attend follow-up visit at the age of 18.5 years during 2012–2014.
Cervical and oropharyngeal samples for HPV DNA testing were obtained on each visit. Setting the age for attending the follow-up visit at 18.5 years was per protocol to allow a minimum of 3 years between vaccination and cervical sampling. During the follow-up visit, the attendees filled-in a questionnaire on living conditions, life-habits and sexual behavior. At age 18.5 years, a cross-vaccination with either the HBV vaccine or the AS04-HPV-16/18 vaccine was offered to the attendees. Female attendees (139), who had moved between Arm C and Arm A or B communities during the follow-up by age 18.5 years were removed from the final analyses.

Laboratory analyses
All samples were analyzed by PCR for HPV DNA. HPV typing was performed by a broad-spectrum PCR SPF10-LiPA25 (Labo Biomedical Products, Rijswijk, the Netherlands) using HPV-specific hybridization probes enabling detection of 14 oncogenic HPV types (HPV-16, 18, 31, 33, 35, 39, 45, 51, 52, 56, 58, 59, 66 and 68) and 11 nononcogenic HPV types (HPV-6, 11, 34, 40, 42, 43, 44, 53, 54, 70 and 74). To ensure maximum sensitivity in the detection of HPV-16, 18, 31, 33, 35, 39, 45, 51, 52, 56, 58, 59, 66, 68, 6 and 11, samples initially considered to be SPF-10/DEIA positive for HPV were re-evaluated by the multiplex type-specific PCR.

Statistical analyses
The number of subjects invited to participate in the study and the number of subjects enrolled was tabulated by gender, for birth cohorts 1994, 1995 and overall.

The analysis of VE was done on female study participants and was based on the total enrolled cohort which included all study participants from all communities, including subjects who only completed the behavioral questionnaire at 18.5 years of age. For the analysis of a specific endpoint, only subjects with measured endpoints were considered.

The statistical analysis of VE of the AS04-HPV-16/18 vaccine against oropharyngeal infection was done by comparing the prevalence in females vaccinated with the AS04-HPV-16/18 vaccine from pooled arms A and B over the prevalence in all females from Arm C. As a post hoc analysis, only point estimated of the VE was provided due to the low number of events. It was computed as 1 minus the prevalence (Supporting Information Table S1) in females vaccinated with the AS04-HPV-16/18 vaccine from pooled arms A and B over the prevalence in all females from Arm C.

Data availability
Anonymized individual participant data and study documents can be requested for further research from www.clinicalstudydatarequest.com.

Results
Recruitment and follow-up visit
Recruitment to the trial was population-based and essentially balanced across all three study arms (Table 1). Also, attendance to the follow-up visit at the age of 18.5 years by vaccinated female trial participants and nonvaccinated female residents of the different trial arm communities was similar.

Vaccine effectiveness against oropharyngeal HPV infections
We found high to moderate VE against oropharyngeal hrHPV infections. VE against vaccine HPV types 16/18 was high (82.4%; 95% CI 47.3, 94.1), and indistinguishable between the

Table 1. Number (%) of invited, enrolled and followed females by vaccination arm and birth cohort in 2008–2009* and 2012–2014 (follow-up visit)

| Arm | Birth cohort | 1994 | 1995 | Total |
|-----|--------------|------|------|-------|
| A   | Invited      | 3,063| 2,846| 5,909 |
|     | Enrolled     | 1,964(64.1)| 1,660(58.3) | 3,624(61.3) |
|     | Followed     | 831  | 775  | 1,606 |
| B   | Invited      | 3,572| 3,408| 6,980 |
|     | Enrolled     | 2,059(57.6)| 1,913(56.1) | 3,972(56.9) |
|     | Followed     | 813  | 773  | 1,586 |
| C   | Invited      | 3,040| 3,027| 6,067 |
|     | Enrolled     | 1,890(62.2)| 1,789(59.1) | 3,679(60.6) |
|     | Followed     | 846  | 833  | 1,679 |
| All | Invited      | 9,675| 9,281| 18,956 |
|     | Enrolled     | 5,913(61.1)| 5,362(57.8)| 11,275(59.5) |
|     | Followed     | 2,490| 2,381| 4,871 |

Arm A: 90% of vaccinated females and males received the AS04-adjuvanted HPV-16/18 (AS04-HPV-16/18) vaccine, 10% received the hepatitis B virus (HBV) vaccine. Arm B: 90% of vaccinated females received the AS04-HPV-16/18 vaccine, 10% of vaccinated females and 100% vaccinated males received the HBV vaccine. Arm C: 100% of vaccinated females and males received the HBV vaccine. The total enrolled cohort included all study participants from all communities, including subjects who only completed the behavioral questionnaire at 18.5 years of age. The followed females included all subjects with human papillomavirus (HPV) DNA Polymerase Chain Reaction result available for the oral gargle sample taken at the follow-up visit and used in the computation of the total vaccine effectiveness. *Q1/2010, for the 1995 birth cohort.
two types (Table 2). VE against cross-protected HPV types 31/45 and 31/33/45 was 75.3% (95% CI: 12.7, 93.0) and 69.9% (95% CI 29.6, 87.1), respectively.

No VE was found against other hrHPV types (Table 2). VE estimate against the most notable low-risk HPV types 6/11 was low (25.8; 95% CI –21.7, 54.8).

**Discussion**

We found for the AS04-HPV-16/18 vaccine high VE against oropharyngeal infections with the vaccine HPV types 16 and 18 and a moderate VE against a prespecified combination of hrHPV types 31/33/45 not included in the vaccine. These results are in line with the reported, respectively high and moderate VEs of the AS04-adjuvanted HPV-16/18 vaccine, 10% vaccinated females and males received the HBV vaccine. Arm C: 100% of vaccinated females and males received the HBV vaccine.

Arm A: 90% of vaccinated females and males received the AS04-adjuvanted HPV-16/18 (AS04-HPV-16/18) vaccine, 10% vaccinated females and males received the HBV vaccine. Arm B: 90% of vaccinated females received the AS04-HPV-16/18 vaccine. Our study provides further evidence that HPV vaccination could probably reduce oral HPV infections which may translate into protection against HPV-related head and neck cancers.

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Table 2. Total vaccine effectiveness against oropharyngeal infection with HPV in AS04-HPV-16/18 vaccinated females (pooled Arms A/B) vs. non-AS04-HPV-16/18 vaccinated females (Arm C) for birth cohorts 1994–1995 vaccinated in 2007–2009 and attending follow-up visits with oral gargle sampling in 2012–2014.

| HPV type | Arms A/B (N = 3,192) Positives n (%) | Arm C (N = 1,679) Positives n (%) | Arms A/B vs. Arm C VE % (95% CI) |
|----------|----------------------------------|----------------------------------|---------------------------------|
| HPV-6/11 | 41 (1.3) 29 (1.7) 25.8 (–21.7, 54.8) | 81.3 (25.8, 95.3) |
| HPV-16  | 6 (0.2) 19 (1.1) 81.3 (25.8, 95.3) |
| HPV-18  | 4 (0.1) 10 (0.6) 78.9 (32.3, 93.4) |
| HPV-16/18 | 9 (0.3) 27 (1.6) 82.4 (47.3, 94.1) |
| HPV-31  | 2 (0.1) 8 (0.5) 86.8 |
| HPV-33  | 6 (0.2) 9 (0.5) 64.9 |
| HPV-35  | 3 (0.1) 3 (0.2) 47.4 |
| HPV-45  | 1 (0.0) 1 (0.1) 47.4 |
| HPV-52  | 14 (0.4) 12 (0.7) 38.6 |
| HPV-31/45 | 3 (0.1) 9 (0.5) 75.3 (12.7, 93.0) |
| HPV-31/33/45 | 9 (0.3) 16 (1.0) 69.9 (29.6, 87.1) |

Arms A/B: 90% of vaccinated females and males received the AS04-adjuvanted HPV-16/18 (AS04-HPV-16/18) vaccine, 10% received the hepatitis B-virus (HBV) vaccine. Arm C: 100% of vaccinated females and males received the HBV vaccine.

1Only point estimated of VE was provided for this post hoc analysis due to the low number of events.

Abbreviations: CI, confidence interval; HPV, human papillomavirus; n, number of subjects with available results; n (%), number (percentage) of subjects reporting an event; VE, total vaccine effectiveness.

Community-randomized trial mimics national vaccination programs as does the female HPV vaccination coverage of approximately 50% in the HPV-vaccination arms. The early age at vaccination was optimal for the vaccine-induced immune response and to ensure the lowest possible proportion of baseline positives. High attendance rates of both originally vaccinated female participants and unvaccinated female residents at the age of 18.5 years retained the population-based nature of our trial. This yielded robust VE estimates from the comparison of HPV-16/18 vaccinated females in combined Arms A/B vs. non-HPV-16/18 vaccinated females in Arm C.

There was a long lag of 4–6 years in the collection of oral gargle samples postvaccination, while the oropharyngeal baseline HPV infection status in the study participants prior to vaccination was unknown. Identification of hrHPV types by highly sensitive PCR in the oral gargle is, however, the most comprehensive means for the identification of oropharyngeal HPV infections.

Whether the identification of hrHPV type(s) in these cross-sectional samples taken at the age of 18.5 years represents persistent infection can therefore not be judged, and must be considered as a limitation of the study. On the other hand, while 6 months and 12 months persistent positivity for the same HPV type has been used to define persistent cervical or other anogenital infections, very little is known about the duration of oropharyngeal infections. In the future, vaccine efficacy against oropharyngeal persistent hrHPV infection will be documented in an ongoing 10-year follow-up study with serial oral gargle sampling.

In conclusion, high VE against oropharyngeal infections with vaccine HPV types 16/18 and moderate VE against infections with nonvaccine HPV types 31/33/45 was observed in young adult females up to 6 years postvaccination with the AS04-HPV-16/18 vaccine. Our study provides further evidence that HPV vaccination could probably reduce oral HPV infections which may translate into protection against HPV-related head and neck cancers.

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References

1. de Martel C, Plummer M, Vignat J, et al. Worldwide burden of cancer attributable to HPV by site, country and HPV type. Int J Cancer 2017;141:664–70.

2. Gillison ML, Koch WM, Capone RB, et al. Evidence for a causal association between human papillomavirus and a subset of head and neck cancer. J Natl Cancer Inst 2000;92:709–20.

3. Mork J, Lie AK, Glattre E, et al. Human papillomavirus infection as a risk factor for squamous cell carcinoma of the head and neck. N Engl J Med 2001;344:1125–31.

4. Smeets SJ, Hesselink AT, Speel EJ, et al. A novel algorithm for reliable detection of human papillomavirus in paraffin embedded head and neck specimens. Int J Cancer 2007;121:2465–72.

5. Haeggblom L, Ramqvist T, Tommasino M, et al. Time to change perspectives on HPV in oropharyngeal cancer. A systematic review of HPV prevalence per oropharyngeal sub-site the last 3 years. Papillomavirus Res 2017;4:1–11.

6. IARC. A review of human carcinogens. Monographs on the evaluation of carcinogenic risks to humans, vol. 100B. Lyon: IARC, 2009.

7. Näsman A, Attnér P, Hammarstedt L, et al. Incidence of human papillomavirus (HPV) positive tonsillar carcinoma in Stockholm, Sweden: an epidemic of viral-induced carcinoma? Int J Cancer 2009;125:362–6.

8. Ramey T, Dalianis T. Oropharyngeal cancer epidemic and human papillomavirus. Emerg Infect Dis 2010;16:1671–7.

9. Cheraghlou S, Yu PK, Otremba MD, et al. Treatment deintensification in human papillomavirus-positive oropharynx cancer: outcomes from the National Cancer Data Base. Cancer 2017;124:717–26.

10. Schroeder L, Wichmann G, Willner M, et al. Antibodies against human papillomaviruses as diagnostic and prognostic biomarker in patients with neck squamous cell carcinoma from unknown primary tumor. Int J Cancer 2017;142:1361–8.

11. Shah SS, Senapati S, Klacsman F, et al. Current technologies and recent developments for screening of HPV-associated cervical and oropharyngeal cancers. Cancers (Basel) 2016;8:885.

12. Beachler DC, Kreimer AR, Schiffman M, et al. Costa Rica HPV vaccine trial (CVT) group. Multi-site HPV16/18 vaccine efficacy against cervical, anal and oral HPV infection. J Natl Cancer Inst 2016;108:djv302.

13. Lupato V, Holzinger D, Höfler D, et al. Prevalence and determinants of oral human papillomavirus infection in 500 young adults from Italy. PLoS One 2017;12:e0170091.

14. Grütt N, Mbuya W, Ternhag A, et al. Human papillomavirus prevalence in mouthwashes of patients undergoing tonsillectomy shows dominance of HPV69, without the corresponding antibodies in the oral cavity. Br J Cancer 2016;114:409–16.

15. Rusan M, Klug TE, Henrissen JI, et al. Prevalence of tonsillar human papillomavirus infections in Denmark. Eur Arch Otorhinolaryngol 2015;272:2505–12.

16. Handsرعا A, Schellenbacher C, Hattel A, et al. Human papillomavirus vaccination induces neutralising antibodies in oral mucosal fluids. Br J Cancer 2016;114:409–16.

17. Pinto LA, Kemp TJ, Torres BN, et al. Quadrivalent human papillomavirus (HPV) vaccine induces HPV-specific antibodies in the oral cavity: results from the mid-adult male vaccine trial. J Infect Dis 2016;214:1276–83.

18. Chaturvedi AK, Graubard BI, Broussard T, et al. Effect of prophylactic human papillomavirus (HPV) vaccination on oral HPV infections among young adults in the United States. J Clin Oncol 2017;36:262–7.

19. Lehtinen M, Apter D, Baussano I, et al. Characteristics of a cluster-randomized, phase IV human papillomavirus vaccination effectiveness trial. Vaccine 2015;33:1284–90.

20. Lehtinen M, Paavonen J, Wheeler CM, et al. Overall efficacy of HPV-16/18 vaccine against the most stringent cervical pre-cancer end-points: end-of-study report of a double blind, randomized trial. Lancet Oncol 2012;13:89–99.

21. Wheeler CM, Castellsague X, Garland SM, et al. Efficacy of the HPV-16/18 AS04-adjuvanted vaccine against cervical infection and precancer caused by non-vaccine oncogenic HPV types (PATRICIA trial). Lancet Oncol 2012;13:100–10.

22. Cameron RL, Kavanagh K, Pan J, et al. Human papillomavirus prevalence and herd immunity after introduction of vaccination program, Scotland, 2009–2013. Emerg Infect Dis 2016;22:56–64.

23. Lehtinen M, Luostarinen T, Vänskä S, et al. Gender-neutral provides improved control of human papillomavirus types 18/31/33/35 through herd immunity: results of a community randomized trial (III). Int J Cancer 2018;143:2299–31.

24. Dona MG, Pichl B, Rollo F, et al. Human papillomavirus detection in matched oral rinses, oropharyngeal and oral brushings of cancer-free high-risk individuals. Oral Oncol 2019;91:1–6.