HPV type concordance in sexual couples determines the effect of condoms on regression of flat penile lesions

We earlier demonstrated, in a randomised clinical trial, that the regression time of flat penile lesions in male sexual partners of women with cervical intraepithelial neoplasia (CIN) was shorter in men who used condoms compared to those who did not. To further evaluate this finding, we examined whether the effect of condom use on the regression of flat penile lesions depends on the presence of human papillomavirus (HPV) type concordance in sexual couples, as determined in cervical and penile scrapes by GP5 +/6 + PCR testing. A Cox model with time-dependent covariates showed a beneficial effect of condoms on regression of flat penile lesions in concordant couples (hazard ratio 2.63, 95% CI 1.07–6.48) but not in those who were nonconcordant. When both partners harboured different HPV types, no effect of condoms was found (hazard ratio 0.90, 95% CI 0.27–2.96). Delayed regression of flat penile lesions was associated with either stable lesions or with new penile lesions developing at sites surrounding pre-existing lesions suggesting reinfection of the penile epithelium. We conclude that condom use blocks sexual HPV transmission by preventing reinfection and development of new penile lesions in men who are susceptible to the same type as present in the female partner.

Keywords: condom use; flat penile lesions; HPV type concordance

Infection with high-risk types of human papillomavirus (HPV) is the major cause of cervical cancer and its precursor lesions (Schiffman et al, 1993; Walboomers et al, 1999; Bosch et al, 2002; Munoz et al, 2003). Human papillomavirus has also been linked to cancers of the vulva, vagina, anus, and penis (IARC 1995; Melbye and Frisch, 1998). Genital HPV types are transmitted sexually and the risk of infection has been associated with numerous parameters involving sexual behaviour (Franco et al, 1995; Bosch et al, 1996; Kjaer et al, 2001).

Partner studies have shown that penile lesions are found in more than half of the male partners of women with cervical intraepithelial neoplasia (CIN) (Barrasso et al, 1987; Campion et al, 1988; Hippelainen M et al, 1991; Bleeker et al, 2002). Flat lesions are reported as the most prevalent lesion type and their relationship with HPV has been established (Barrasso et al, 1987; Campion et al, 1988; Hippelainen M et al, 1991; Hippelainen MI et al, 1993b; Bleeker et al, 2002). Since flat penile lesions are only well visible after acetowhiteing, their presence remains often unnoticed. This explains why the natural history of penile lesions and HPV infection in men has not been studied extensively and is largely unknown.

We have recently shown that the presence of penile HPV is associated with delayed regression whereas condom use revealed accelerated regression (Bleeker et al, 2003). The beneficial effect of condom intervention on regression of penile lesions suggests that a sexual transmittable factor, that is, most likely cervical HPV, influences the clinical course of these lesions. To evaluate this hypothesis, the effect of condom use on regression of flat penile lesions was studied in relation to the presence of HPV type concordance between sexual partners. Since use of condoms and presence of concordance could change during this prospective study, effects of condom use were estimated by a Cox model with time-dependent covariates.

**PATIENTS AND METHODS**

**Study population**

The study population consisted of men recruited at the colposcopy clinic of the Albert Schweitzer Hospital, Dordrecht, the Netherlands, from January 1995 to June 2002. These men participated in a randomised clinical trial, studying the influence of condom use on regression of CIN and penile lesions in sexual couples. Details of
this trial have been described recently (Bleeker et al, 2003; Hogewoning et al, 2003). Men were screened for penile lesions by penoscopy and a penile scrape was taken for HPV PCR testing. Women were screened by colposcopy and scrapes were taken for cytology and HPV testing. Baseline cervical biopsy specimens were taken for histopathological evaluation. Couples were randomised to either condom use or not using a block randomisation scheme with block size two. Condoms were proposed for at least 3 months and randomisation occurred independent of the presence of HPV, penile lesions, grade of CIN lesion, or surgical treatment of the CIN lesion. Exclusion criterion was regular condom use at baseline. Follow-up examinations were carried out after 3 and 6 months and subsequently every 6 months. In the condom arm, condom use was verified at each visit by asking the frequency of failures to use condoms. Partners were not asked separately for the number of failures and, in case of discrepancies between partners, the couples themselves made a consensus. Also, couples assigned to the noncondom arm were asked during each follow-up visit whether they have used condoms but none of them reported condom use. To obtain information on lifestyle habits, including sexual behaviour, couples were asked to complete a questionnaire. This questionnaire was introduced in 1999 and completed, independently by each partner, in separate rooms. The medical ethics review board of the hospital approved the study protocol (protocol number: CGE/95/238) and the participants signed informed consents.

Data collection

Penoscopic examination Penoscopic examination and specimen collection were performed as described earlier by us (Bleeker et al, 2002). Briefly, using a Cervex brush™, a penile scrape for HPV testing was taken to collect cells from the glans, corona, sulcus, frenulum, and inner part of the foreskin. Subsequently, a colposcope was used to identify flat penile lesions after the application of 3% acetic acid solution. Follow-up penoscopic examinations were performed blinded of data from previous visits. Findings were written on a standardised form and penile lesions were documented by digital photography. Photographs were subsequently reviewed and graded by an experienced dermatologist (TMS), unaware of any clinical data. In case of discrepancies (<5%) between the initial finding and the review result, a consensus diagnosis was made with the gynaecologist who made the photographs and the dermatologist who reviewed the photographs.

Human papillomavirus testing Processing of specimens and testing for HPV was carried out at the Department of Pathology, VU University Medical Center, as described previously (Jacobs et al, 1997; Bleeker et al, 2002). In short, DNA quality was tested by β-globin PCR and testing for HPV was carried out using the HPV GP5+/6+ PCR enzyme immunoassay (HPV PCR-EIA), to identify 14 high-risk (16, 18, 31, 33, 35, 39, 45, 51, 52, 56, 58, 59, 66, and 68) and six low-risk (6, 11, 40, 42, 43, and 44) HPV genotypes with a high-risk and a low-risk HPV probe cocktail, respectively. Additionally, individual typing was performed for HPV 16, 18, 31, 33, 6, and 11, using type-specific probes in an EIA format. Samples, which were inadequate for HPV testing, that is, were negative in both β-globin and HPV PCR assays, were excluded from the analysis.

Regression of flat penile lesions

In 75% of the men with flat penile lesions, more than 5 mm² of the penile epithelium was affected and in 94% of them more than one lesion was present (Bleeker et al, 2005). Follow-up of individual flat penile lesions was sometimes difficult, especially when lesions were small and present on multiple sites of the penis. Larger lesions were usually localized on the same site of the penile epithelium and could vary in size. Owing to this diverse clinical presentation, regression was defined as disappearance of all detectable flat lesions, that is, no flat lesions at penoscopy.

Statistical analysis

Differences in characteristics of the condom and the noncondom groups were assessed by use of χ² or Student’s t-tests (two-tailed, P < 0.05), where appropriate.

The outcome of interest in this study was regression of flat penile lesions. To study the complex interrelationship between HPV presence, concordance, and condom use on the regression of the penile lesions, we choose an approach in which changes in condom use or presence of female HPV infection during the study could be included. Therefore, to examine whether the effect of condom use on regression of flat penile lesions was related to the presence of HPV type concordance in sexual couples, Cox regression analysis with time-dependent covariates (condom use and presence of female HPV) was performed to compute hazard ratios and 95% confidence intervals (CI). In the modelling process, hazard ratios for the effect of condom use were computed using the noncondom group as the reference group in couples with a comparable HPV status.

The HPV status in men was defined as positive or negative when penile scrapes were ever or never HPV positive during the study, respectively (Bleeker et al, 2003). Other determinants assessed were CIN grade of the female partner at baseline histology (either ≤CIN 1 or ≥CIN 2) and age of the men when entering the study.

Originally, sexual couples were considered as HPV type concordant when both sexual partners were positive for HPV 16, 18, 31, 33, 6, 11, other high-risk HPVs (not HPV type 16, 18, 31, or 33) and/or other low-risk HPV (not HPV type 6 and 11) during the study. This definition was used because of practical reasons that individual typing was only performed for HPV 16, 18, 31, 33, 6, and 11, whereas other high-risk HPVs and other low-risk HPVs each typed as a group. A more restrictive definition of concordance was subsequently used including only the couples that were concordant for 16, 18, 31, 33, 6, and/or 11 and excluding the couples that were concordant for other high-risk and/or other low-risk HPV types.

For Cox regression analyses with time-dependent covariates, Stata 6.0 was used and for the remaining analyses SPSS version 9.0 software.

RESULTS

Study population

A total of 139 men had flat penile lesions at baseline. Six of them used condoms at baseline and were excluded from this study. No follow-up results were available from 14 men, and in one female partner the presence of HPV could not be determined due to inadequate sample quality at baseline. The remaining 118 men formed the study population. At baseline, 61 (51.7%) couples started condom use and 57 (48.3%) did not. The median duration of condom use was 5.3 months (range 2.5–53.7). Failures of condom use were reported by six couples with a median of 1.5 times during the follow-up period (range 1–2). The median follow-up time of the study population was 12.5 months (range 2.5–77.5): 12.9 months (range 3.0–68.5) in the noncondom and 12.3 months (range 2.5–77.5) in the condom group (P = 0.388).

Baseline characteristics of the study population are presented in Table 1 and no statistically significant differences were found between the condom and the noncondom group. The mean age of men was 38.7 years (range 28.1–57.7 years) and of women 36.2 years (range 22.6–54.9). Representative biopsy specimens were
available for 111 women: 17 had no CIN (15.3%), 34 CIN 1 (30.6%), 39 CIN 2 (35.1%), and 21 ≥ CIN 3 (18.9%). At baseline, 85.6% of the women and 68.6% of the men were positive for HPV DNA. Human papillomavirus 16 was found in 59.4 and 55.9% of the positive women and men, respectively, and their prevalences were not statistically different in the condom and the noncondom group ($P = 0.496$ in women and $P = 0.775$ in men). Women with HPV at baseline ($n = 101$) had a median duration of HPV infection of 9.2 months and were not statistically different in the condom and noncondom group ($P = 0.852$).

During the study, at least one concordant HPV type was found in 57 couples: 32 couples of the condom and 25 couples of the noncondom group ($P = 0.350$). Human papillomavirus type concordance was found for HPV 16 in 59.4 and 55.9% of the positive women and men, respectively, and their prevalences were not statistically different in the condom and the noncondom group ($P = 0.496$ in women and $P = 0.775$ in men). Women with HPV at baseline ($n = 101$) had a median duration of HPV infection of 9.2 months and were not statistically different in the condom and noncondom group ($P = 0.852$).

**Table 1** Baseline characteristics of the study population

|                      | Condom− | %       | N       | %       | Condom+ | %       |
|----------------------|---------|---------|---------|---------|---------|---------|
| Subjects             | 118     | 100     | 57      | 48.3    | 61      | 51.7    |
| Age                  |         |         |         |         |         |         |
| Men                  | 38.7    | 38.9    | 3.8     | 36.2    | 36.1    | 36.3    |
| Women                | 36.2    | 36.1    |         | 36.3    |         |         |
| Histological CIN grade in female partner |         |         |         |         |         |         |
| No (representative) biopsy | 7     | 4       | 3       | 7       | 4       |
| No CIN               | 17      | 15.3    | 10      | 18.9    | 7       | 12.1    |
| CIN 1                | 34      | 30.6    | 15      | 28.3    | 19      | 32.8    |
| CIN 2                | 39      | 35.1    | 15      | 28.3    | 24      | 41.4    |
| ≥ CIN 3              | 21      | 18.9    | 13      | 24.5    | 8       | 13.8    |
| Cervical HPV         |         |         |         |         |         |         |
| Negative             | 17      | 14.4    | 11      | 19.3    | 6       | 9.8     |
| Positive             | 101     | 85.6    | 46      | 80.7    | 55      | 90.2    |
| Penile HPV           |         |         |         |         |         |         |
| Inadequate*          | 32      | 19      | 13      |         |         |         |
| Negative             | 27      | 31.4    | 12      | 31.6    | 15      | 31.3    |
| Positive             | 59      | 68.6    | 26      | 68.4    | 33      | 68.8    |

*Could not be determined because samples were inadequate for HPV testing. HPV = human papillomavirus; CIN = cervical intraepithelial neoplasia.

**Table 2** Effect of condom use on regression of flat penile lesions

| Cervical HPV | Penile HPV | Effect of condom use* |
|--------------|------------|-----------------------|
| Nonconcordant| Nonconcordant| Hazard ratio | 95% CI  | $P$  |
| Negative     | Negative   | 1.02                  | 0.10 – 10.11 | 0.986 |
|              | Positive   | 1.04                  | 0.07 – 15.55 | 0.978 |
| Positive     | Negative   | 2.40                  | 0.47 – 12.36 | 0.294 |
|              | Positive   | 0.90                  | 0.27 – 2.96  | 0.862 |
| Concordant   | Concordant | 2.63                  | 1.07 – 6.48  | 0.035 |
| Positive     | Positive   | 2.63                  | 1.07 – 6.48  | 0.035 |

*Cox regression analyses with time-dependent predictors (cervical HPV and condoms) were performed to compute 95% CI of the hazard ratios for the effect of condom use vs no use of condoms. Boldface numbers indicate statistical significance. CI = confidence interval; HPV = human papillomavirus.

**Effect of condom use on regression of flat penile lesions in concordant and nonconcordant sexual couples**

Using a Cox model with time-dependent covariates, we examined whether the effect of condom use on regression of flat penile lesions was related to the presence of HPV type concordance in sexual partners. This analysis showed that the beneficial effect of condoms on regression of flat penile lesions was present in concordant couples (hazard ratio 2.63, 95% CI 1.07 – 6.48, $P = 0.035$, Table 2) but not in those who were nonconcordant. Also, no effect of condoms was found when both partners were positive but nonconcordant (hazard ratio 0.90, 95% CI 0.27 – 2.96, $P = 0.861$).

When we restricted the analysis to couples concordant for HPV types 6, 11, 16, 18, 31, and 33, which were individually typed during this study, similar effects of condom use were obtained.

Regression of flat penile lesions was not related to the age of the men (hazard ratio 0.99, 95% CI 0.95 – 1.03, $P = 0.501$) or the histological CIN grade (hazard ratio 1.27, 95% CI 0.71 – 2.25, $P = 0.401$).
DISCUSSION

Our data, collected in a randomised clinical trial, earlier demonstrated that the regression time of flat penile lesions was shorter in men who used condoms compared to those who did not (Bleeker et al., 2003). We hypothesised that sexually transmittable HPV of the women delayed regression of flat penile lesions in their male partner and that condoms could effectively block HPV transmission. In the present study, we examined whether the effect of condom use on regression of flat penile lesions depended on the presence of HPV concordance between sexual partners. Indeed, using a Cox model in which changes on condom use and presence of HPV concordance during this study were included, a beneficial effect of condoms on regression of penile lesions was found for HPV type concordant couples, but not for those who were not concordant.

The presence of penile lesions associated with a given HPV type indicates that the man is susceptible for this particular HPV type. In men with penile lesions who harboured a similar HPV type as their female partner, we observed a delayed regression when couples did not use condoms compared to those who did. This finding suggests that failure of penile lesions to cure depends on reinfection by cervical HPV from the female partner for which the man is apparently susceptible. This is substantiated by the observation that new penile lesions often developed next to pre-existing ones. The beneficial effect of condom use in HPV concordant couples can be explained by blocking HPV transmission during sexual intercourse, thereby preventing reinfection and development of new penile lesions. In nonconcordant couples, we found no effect of condom use on regression of penile lesions. Apparently, these men are not susceptible for the development of new lesions by the HPV type, which is present in the female partner. Consequently, the clinical implication of our findings would be that condoms sort only effect in heterosexual couples that are positive for the same HPV type and that, in these couples, condom use should be continued as long as the female partner is HPV positive. In addition, condom use could also be advocated in sexual couples of whom the woman is HPV positive and the male HPV negative. Another issue is that we did not ask the couples to keep a log of their condom use. Therefore, because of potential false negativity in penile scrapes, the currently defined group of nonconcordant couples may contain some concordant couples as well. This may explain the hazard ratio of 2.40 (95% CI 0.47–12.36) in nonconcordant couples in whom the woman was HPV positive and the man negative. Another issue is that we did not ask the couples to keep a log of their condom use. Therefore, recall bias cannot be completely excluded but it is unlikely that this substantially influenced the data as the number of failures to use condoms was low. Moreover, failures to use condoms would most likely lower the positive effect of condoms on regression of the penile lesions, meaning that the effect of condoms would even be more beneficial when there were no failures of condom use at all.

In conclusion, the beneficial effect of condoms on regression of flat penile lesions was only found when sexual couples were HPV concordant. The effect of condoms is probably the result of blocking transmission of female HPV to the penis, thereby preventing reinfection and induction of a penile lesion to which a man is susceptible. As HPV testing becomes more widely used for both the management of women with HPV-related disorders and in cervical screening programs, accurate information on how to protect against HPV is needed. Our data suggest that condom use can reduce the spread of HPV in a sexually active population. Our findings warrant further studies on this topic.

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So far, studies on the effectiveness of condoms in relation to the prevention of HPV infection and related disorders were mainly focused on women and available data were too inconsistent to provide precise estimates (Manhart and Koutsky, 2002). Only a few studies were performed in men and reported that consistent condom use reduced the risk for genital warts or subclinical HPV infection (Hippelainen M et al., 1993a; Wen et al., 1999). Of interest is the work of Hippelainen MI et al (1994) who found that condom use had a positive effect on the cure rate of genital HPV lesions in male partners of HPV-infected women. However, all studies evaluating condom use were either cross-sectional or observational and not explicitly designed to evaluate condom use prospectively.

In this partner study, however, it should be realised that grouping into concordant and nonconcordant may be subject to some degree of underestimation of concordant sexual couples. Nondetection of HPV in penile scrapes does not necessarily reflect absence of HPV (Bleeker et al., 2003). Compared to cervical scrapings, low cell numbers are collected by scraping the penile epithelium and consequently low numbers of infected cells might have been missed during the sampling. Therefore, because of potential false negativity in penile scrapes, the currently defined group of nonconcordant couples may contain some concordant couples as well. This may explain the hazard ratio of 2.40 (95% CI 0.47–12.36) in nonconcordant couples in whom the woman was HPV positive and the man negative. Another issue is that we did not ask the couples to keep a log of their condom use. Therefore, recall bias cannot be completely excluded but it is unlikely that this substantially influenced the data as the number of failures to use condoms was low. Moreover, failures to use condoms would most likely lower the positive effect of condoms on regression of the penile lesions, meaning that the effect of condoms would even be more beneficial when there were no failures of condom use at all.

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