



Prevalence of High-Risk Human Papillomavirus Infections before and after Cervical Lesion Treatment, among Women Living with HIV

Timothée Dub, Sophie Le Coeur, Nicole Ngo-Giang-Huong, Wanmanee Matanasarawut, Pornnapa Suriyachai, Kannikar Saisawat, Chaiwat Putiyanun, Sudanee Buranabanasatean, Prattana Leenasirimakul, Samreung Randaeng, et al.

► To cite this version:

Timothée Dub, Sophie Le Coeur, Nicole Ngo-Giang-Huong, Wanmanee Matanasarawut, Pornnapa Suriyachai, et al.. Prevalence of High-Risk Human Papillomavirus Infections before and after Cervical Lesion Treatment, among Women Living with HIV. Journal of Clinical Medicine, 2021, 10 (14), pp.3133. 10.3390/jcm10143133 . hal-03510072

HAL Id: hal-03510072

<https://hal.science/hal-03510072>

Submitted on 4 Jan 2022

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.





L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Distributed under a Creative Commons Attribution 4.0 International License

Article

Prevalence of High-Risk Human Papillomavirus Infections before and after Cervical Lesion Treatment, among Women Living with HIV

Timothée Dub ^{1,2,*} , Sophie Le Cœur ^{2,3} , Nicole Ngo-Giang-Huong ^{2,4} , Wanmanee Matanasarawat ⁵, Pornnapa Suriyachai ⁶, Kannikar Saisawat ⁷, Chaiwat Putiyanun ⁸, Sudanee Buranabansatean ⁹, Prattana Leenasirimakul ¹⁰, Samreung Randaeng ¹¹ and Tristan Delory ^{2,3,4,12} 

- ¹ Department of Health Security, Finnish Institute for Health and Welfare, 00160 Helsinki, Finland
 - ² Faculty of Associated Medical Sciences, Chiang Mai University, Chiang Mai 50200, Thailand; lecoeur@ined.fr (S.L.C.); Nicole.Ngo-Giang-Huong@phpt.org (N.N.-G.-H.); delory.tristan@gmail.com (T.D.)
 - ³ Institut National d'Études Démographiques (INED), 93322 Aubervilliers, France
 - ⁴ Institut de Recherche pour le Développement (IRD) UMI 174-PHPT, 13002 Marseille, France
 - ⁵ Ministry of Public Health, Lamphun Hospital, Lamphun 51000, Thailand; ogyn.lph@gmail.com
 - ⁶ Ministry of Public Health, Phayao Provincial Hospital, Phayao 56000, Thailand; whan76_1@hotmail.com
 - ⁷ Ministry of Public Health, Chiangrai Prachanukroh Hospital, Chiang Rai 57000, Thailand; kannikarsai@gmail.com
 - ⁸ Ministry of Public Health, Chiang Kham Hospital, Chiang Kham 56110, Thailand; putiyanun@hotmail.com
 - ⁹ Ministry of Public Health, Mae Chan Hospital, Chiang Rai 57110, Thailand; sudanee2011@windowslive.com
 - ¹⁰ Ministry of Public Health, Nakornping Hospital, Chiang Mai 50180, Thailand; leeprattana@gmail.com
 - ¹¹ Faculty of Medicine, Department of Pathology, Chiang Mai University, Chiang Mai 50200, Thailand; srangdaee@mail.medicine.cmu.ac.th
 - ¹² Délégation à la Recherche Clinique et à l'Innovation (DRCI), Centre Hospitalier Annecy-Genevois, 74370 Epargny Mets-Tessy, France
- * Correspondence: timothee.dub@thl.fi; Tel.: +358-29-524-6177



Citation: Dub, T.; Le Cœur, S.; Ngo-Giang-Huong, N.; Matanasarawat, W.; Suriyachai, P.; Saisawat, K.; Putiyanun, C.; Buranabansatean, S.; Leenasirimakul, P.; Randaeng, S.; et al. Prevalence of High-Risk Human Papillomavirus Infections before and after Cervical Lesion Treatment, among Women Living with HIV. *J. Clin. Med.* **2021**, *10*, 3133. <https://doi.org/10.3390/jcm10143133>

Academic Editors: Aldo Venuti and Jose Bordon

Received: 2 April 2021

Accepted: 15 July 2021

Published: 15 July 2021

Publisher's Note: MDPI stays neutral with regard to jurisdictional claims in published maps and institutional affiliations.



Copyright: © 2021 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (<https://creativecommons.org/licenses/by/4.0/>).

Abstract: Even when receiving combination antiretroviral therapy, women living with HIV are at high risk of human papillomavirus (HPV) infection and/or cervical lesions, including cancer. Using data from the PapilloV prospective cohort, we evaluated the prevalence of high-risk HPV (HR-HPV) infections after cervical lesion treatment and investigated factors associated with their carriage. Women were followed up for three years with annual Pap smear and HPV genotyping. We offered treatment to women presenting either a Pap smear with high-grade squamous intraepithelial lesion or higher, and/or a biopsy showing cervical intraepithelial neoplasia II or III. We compared the prevalence of HR-HPV infection at the time of first treatment indication and at the end of follow-up among women who received treatment and those who did not. Overall, 46 women had treatment indication. HR-HPV prevalence significantly decreased from 67% to 27% (p value = 0.001) in the 30 women who received treatment, while it did not significantly decrease (from 56% to 38%) in the 16 women who did not (p value = 0.257). Due to lack of statistical power, the 40% relative difference in HR-HPV carriage between treated and untreated women was not significant. In women living with HIV, the treatment of a cervical lesion may be beneficial for clearing HR-HPV infections.

Keywords: human papillomavirus; women living with HIV; cervical lesion

1. Introduction

Women living with HIV are at increased risk of presenting with a human papillomavirus (HPV) infection and/or cervical lesions, including cancer, even when receiving combination antiretroviral therapy [1–3].

In Thailand, among women living with HIV, the prevalence of cervical infection ranges from 25% to 62% for all types of HPV, and from 19% to 43% for high-risk HPV (HR-HPV) [4–8].

In a study evaluating the effect of cervical lesion treatment on HR-HPV carriage, women who had undergone punch biopsies had faster clearance of HR-HPV infections compared to the natural course of infection [9]. After treatment for cervical lesions, more women living with HIV had a persistent HR-HPV infection, compared to HIV-negative women [10]. In a larger study conducted in Nigeria, regardless of treatment, clearance of HR-HPV infection was similar (~8%) among HIV-negative and -positive women [11]. Independently of treatment, younger age was associated with faster HR-HPV clearance [12,13].

Among women living with HIV with a treatment indication for cervical lesion, we evaluated the impact of treatment on HPV carriage, including HR-HPV, between the time of cervical lesion detection and the end of follow-up, by comparing women who received treatment to those who did not. We also investigated potential factors associated with persistent carriage of HR-HPV.

We used data from the PapilloV cohort study, in which 829 women living with HIV in Thailand were screened annually for HPV infection and cervical lesions over three years.

2. Materials and Methods

The PapilloV study was a prospective, multicenter cohort of women living with HIV, receiving antiretroviral therapy, nested in a larger HIV cohort in Thailand [14]. Women were recruited across 24 hospitals in Thailand. The study procedure has been detailed elsewhere [4].

In short, women were followed up for three years. Annual visits comprised a gynecological exam, including a Pap smear and HPV testing with genotyping. Colposcopy was performed if an abnormal Pap smear showed atypical squamous cells of undetermined significance (ASC-US) or higher-grade lesions and/or presence of either HR-HPV or potential HR-HPV (pHR-HPV) infections. Biopsy was performed in case of visible lesion at colposcopy.

Cervical lesion treatment was indicated in case of a Pap smear showing high-grade squamous intraepithelial lesion (HSIL) or higher and/or in case of a biopsy showing cervical intraepithelial neoplasia (CIN) II or III. Women underwent treatment as per physician decision. The study investigation team had no influence on whether treatment was offered or not to the study participants.

HPV genotyping of cervical specimens was conducted using HPV-PapilloCheck (Greiner Bio-One, Frickenhausen, Germany). Genotypes considered high-risk HPV were the following: HPV 16, 18, 31, 33, 35, 39, 45, 51, 52, 56, 58, 59, and 68, while HPV 6, 11, 40, 42, 43, and 44/55 were considered as low-risk HPV genotypes. Due to the limited evidence regarding their potential carcinogenicity, the following genotypes (HPV 53, 66, 70, 73, and 82) were classified as potentially high-risk genotypes (pHR-HPV) [4]. Clinicians deciding for treatment indication were not made aware of the detailed HPV genotyping results but only of the presence of HR- or pHR-HPV.

The study protocol was approved by the ethics committees at the Institute for the Development of Human Research Protections, Ministry of Public Health, Thailand; the Faculty of Associated Medical Sciences, Chiang Mai University, Thailand; and at local hospitals, if established. The PapilloV study was registered at clinicaltrials.gov (NCT01792973).

In this analysis, we included women with cervical treatment indication, either at baseline or anytime during follow-up. We compared their characteristics at the time of treatment indication using the chi-squared test or the Fisher exact test (when required) for categorical variables and the Mann–Whitney U test for numerical values.

We compared the prevalence of each type of HPV infection (HR, pHR, and LR) at the time of the first treatment indication and at the end of follow-up among women who received treatment and among those who did not. Prevalence was compared using McNemar's tests for paired nominal data (before vs. after in the same woman).

We investigated the possible characteristics associated with the presence of an HR-HPV infection by the end of the follow-up period, using bivariate logistic and exact logistic regressions adjusted on having received treatment. We selected the following variables a

priori: age, CD4 cell count, HIV load, delay between ART initiation and cervical treatment indication, Nadir CD4 cell count, and cervical cytology at the end of the follow-up. Two time points were used in estimating prevalence: the time of treatment indication and at the end of follow-up. Hence, to be included, women had to have at least two HPV samples collected.

3. Results

From February 2012 to June 2013, 829 women were recruited in the cohort. Baseline characteristics, prevalence of HPV infections, and cervical lesions have been described elsewhere (4).

Over the median follow-up of 34.9 months (32.0 to 36.8) and the 2073 women-months of follow-up, 49 women had cervical lesions fulfilling treatment indication. Three of them were excluded from the analysis because of missing HPV samples either before or after treatment. The median duration between the initial and final HPV sample was 24.4 months [19.5 to 35.1]. Figure 1 describes the flow chart of women selection for this analysis.

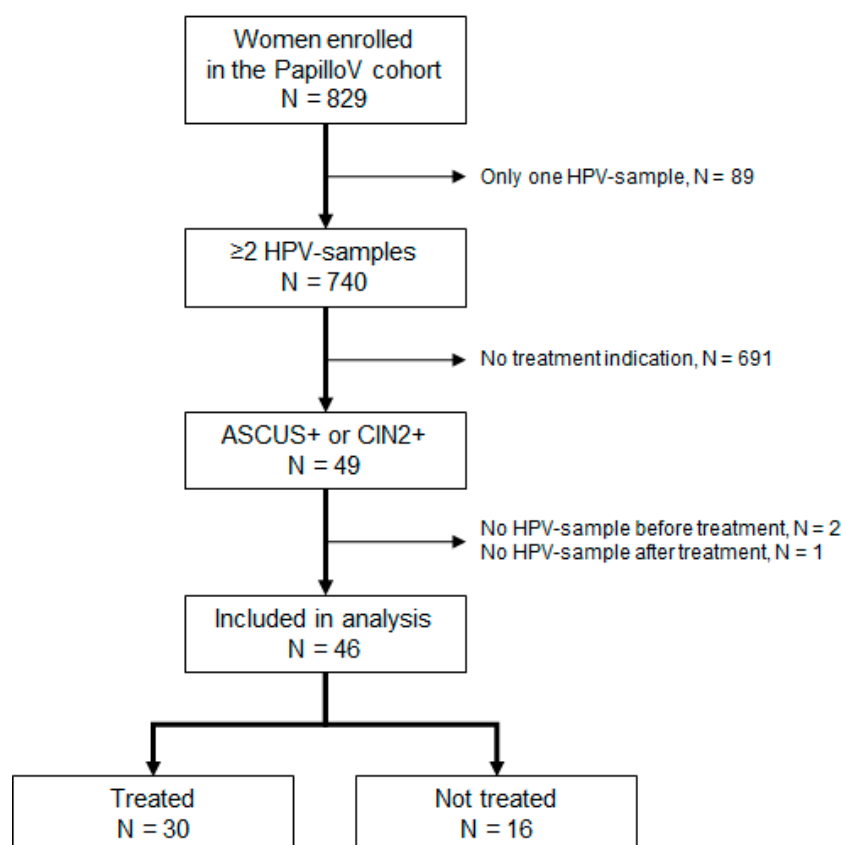


Figure 1. Flow chart of patient selection.

Among the 46 women included in the analysis, 14 (30%) had an HSIL on Pap smear, 12 (26%) had a CIN2, and 20 (43%) a CIN3 on the biopsy at the time of treatment indication. Thirty women received cervical treatment while 16 did not. Out of 30 women treated, 24 were treated once: among them, 22 experienced Loop Electrosurgical Excision Procedure (LEEP) and one participant a cold knife conization, while information on treatment type was missing for one participant. Among the six women who received two successive treatments: two had benefited from LEEP twice, while information was incomplete for four women. Apart from Pap smear and biopsy results, treated and untreated women had similar characteristics at the time of treatment indication (Table 1).

Table 1. Baseline characteristics at the time of indication among women who received treatment and women who did not.

	Received Treatment (<i>n</i> = 30)	Did Not Receive Treatment (<i>n</i> = 16)	<i>p</i> Value
	n (%), Median (IQR)	n (%), Median (IQR)	
Age (years) *	39.5 (33.1–43.7)	39.4 (35.4–42.5)	0.890
CD4 cell count			
>250 cells/mm ³	29 (97)	15 (94)	1.000
<250 cells/mm ³	1 (3)	1 (6)	
HIV load			
<1.7 log10 copies/mL	27 (90)	15 (94)	1.000
>1.7 log10 copies/mL	3 (10)	1 (6)	
Time since ART initiation (years) *	2.2 (1.0–4.8)	3.2 (2.2–4.4)	0.181
HPV infections			
HR-HPV infections	20 (67)	9 (56)	0.534
pHR-HPV infections	3 (10)	3 (19)	0.405
LR-HPV infections	3 (10)	2 (13)	1.000
Any HPV infection	23 (77)	11 (69)	0.726
Multiple HPV infection	6 (20)	4 (25)	0.720
Biopsy results			
CIN 2 –	1 (3)	8 (73)	<0.001
CIN 2 +	29 (97)	3 (27)	
Pap smear results			
HSIL –	20 (69)	3 (19)	0.002
HSIL +	9 (31)	13 (81)	
Combination of biopsy and Pap smear results			
HSIL +/CIN2 +	8 (28)	0	<0.001
HSIL –/CIN2 +	20 (69)	3 (27)	
HSIL +/CIN2 –	1 (3)	8 (73)	
HSIL –/CIN2 –	0	0	
Duration (months) between treatment indication and the last visit *	24.4 (14.0–34.6]	26.7 (20.5–35.1)	0.356

* Median (IQR), Mann–Whitney test.

At the time of treatment indication, overall, 63% (29/46) of women had at least one HR-HPV infection. The most frequent HR-HPV genotypes were HPV 16 in 10 women (22%) and HPV 52 in seven women (15%), while only three women (7%) had an HPV-18 infection (Tables 2 and 3).

Table 2. Prevalence of HPV infection at the time of indication and by the end of the follow-up, among women who received treatment for a cervical lesion.

		Before Treatment Indication	End of Follow-Up	<i>p</i> Value
Genotypes		(<i>n</i> = 30)		
HIGH-RISK HPV INFECTIONS	HPV16	6 (20%)	1 (3%)	0.059
	HPV 18	1 (3%)	0 (0%)	0.317
	HPV 31	3 (10%)	0 (0%)	0.083
	HPV 33	3 (10%)	1 (3%)	0.157
	HPV 35	0	0 (0%)	-
	HPV 39	3 (10%)	0 (0%)	0.083
	HPV 45	1 (3%)	1 (3%)	1.000
	HPV 51	1 (3%)	0 (0%)	0.317
	HPV 52	5 (17%)	4 (13%)	0.564
	HPV 56	1 (3%)	1 (3%)	1.000
	HPV 58	1 (3%)	0 (0%)	0.317
	HPV 59	1 (3%)	0 (0%)	0.317
	HPV 68	2 (7%)	0 (0%)	0.157
HR-HPV regrouped		20 (67%)	8 (28%)	0.001
POTENTIALLY HIGH-RISK HPV INFECTIONS	HPV 53	1 (3%)	0 (0%)	0.317
	HPV 66	1 (3%)	0 (0%)	0.317
	HPV 70	1 (3%)	1 (3%)	1.000
	HPV 73	0 (0%)	0 (0%)	-
	HPV 82	0 (0%)	0 (0%)	-
	pHR-HPV regrouped	3 (10%)	1 (3%)	0.317
LOW-RISK HPV INFECTIONS	HPV 6	0 (0%)	0 (0%)	-
	HPV 11	0 (0%)	0 (0%)	-
	HPV 40	0 (0%)	0 (0%)	-
	HPV 42	1 (3%)	0 (0%)	0.317
	HPV 43	0 (0%)	0 (0%)	-
	HPV 44/55	2 (7%)	0 (0%)	0.157
	LR-HPV regrouped	3 (10%)	0 (0%)	0.083
Any HPV infection		23 (77%)	9 (30%)	<0.001

Table 3. Prevalence of HPV infection at the time of indication and by the end of the follow-up among women who did not receive treatment for cervical lesion.

		Before Treatment Indication	End of Follow-Up	p Value
Genotypes		(n = 16)		
HIGH-RISK HPV INFECTIONS	HPV16	4 (25%)	1 (6%)	0.083
	HPV 18	2 (13%)	1 (6%)	0.317
	HPV 31	1 (6%)	1 (6%)	-
	HPV 33	1 (6%)	1 (6%)	-
	HPV 35	0 (0%)	0 (0%)	-
	HPV 39	0 (0%)	0 (0%)	-
	HPV 45	0 (0%)	0 (0%)	-
	HPV 51	1 (6%)	2 (13%)	0.317
	HPV 52	2 (13%)	2 (13%)	1.000
	HPV 56	0 (0%)	0 (0%)	-
	HPV 58	0 (0%)	1 (6%)	0.317
	HPV 59	0 (0%)	0 (0%)	-
	HPV 68	0 (0%)	1 (6%)	0.317
	HR-HPV regrouped	9 (56%)	6 (38%)	0.257
POTENTIALLY HIGH-RISK HPV INFECTIONS	HPV 53	1 (6%)	0 (0%)	0.317
	HPV 66	0 (0%)	0 (0%)	-
	HPV 70	1 (6%)	1 (6%)	1.000
	HPV 73	1 (6%)	0 (0%)	0.317
	HPV 82	0 (0%)	0 (0%)	-
	pHR-HPV regrouped	3 (19%)	1 (6%)	0.157
LOW-RISK HPV INFECTIONS	HPV 6	0 (0%)	0 (0%)	-
	HPV 11	2 (13%)	1 (6%)	0.317
	HPV 40	0 (0%)	0 (0%)	-
	HPV 42	0 (0%)	0 (0%)	-
	HPV 43	1 (6%)	0 (0%)	0.317
	HPV 44/55	1 (6%)	1 (6%)	-
	LR-HPV regrouped	2 (13%)	2 (13%)	-
Any HPV infection		11 (69)	6 (38%)	0.059

Among women who received treatment, the prevalence of HR-HPV infection was 67% (IC_{95%}, 48% to 82%) at the time of treatment indication and 27% (IC_{95%}, 13% to 46%) at the end of follow-up, a 60% relative decrease of HR-HPV infection prevalence compared to the baseline (p value = 0.001) (Table 2). Among women who did not receive treatment, the HR-HPV prevalence was 56% (IC_{95%}, 20% to 70%) at baseline versus 38% (IC_{95%}, 16% to 65%) at the end of follow-up, a 32% relative decrease, which was not statistically significant (p value = 0.257) (Table 3).

There was no significant association between prevalence of HR-HPV infection by the end of the follow-up and the fact that women had received treatment (OR = 0.6 (IC₉₅, 0.2 to 2.2), p value = 0.449).

Four women still had cervical lesions at the end of follow-up, two among those who received treatment (2 HSIL) (Appendix A Table A1) and two among those who did not (1 HSIL and 1 CIN3) (Appendix A Table A2). Details of HPV genotypes carriage by the end of follow-up are provided in Table 2 for treated women and in Table 3 for those untreated.

When adjusted for having received treatment, none of the women's characteristics at the time of indication (age, CD4 cell count, HIV-RNA viral load, and antiretroviral therapy duration) were associated with HR-HPV carriage at the end of follow-up (Appendix A Table A3). However, the presence of Pap smear showing ASC-US+ at the end of the follow-up was significantly associated with HR-HPV carriage: all HPV negative women had a normal Pap smear at the end of the follow-up (p value < 0.001).

4. Discussion

We found a significant decrease in HR-HPV carriage among women living with HIV who had been treated for a cervical lesion, while it was not significant among women who had not received treatment. However, the relative difference in decrease of HR-HPV infection prevalence was not significant between the two groups.

Interestingly, after adjusting for treatment, there was no significant association between women's HIV characteristics and the presence of an HR-HPV infection at the end of the follow-up. However, the persistence of abnormal cytology findings (ASC-US+) was correlated with HR-HPV carriage by the end of the follow-up.

Some studies have assessed how treatment of cervical lesions might influence HPV clearance, but few have focused on women well-controlled for HIV infection. Also, it should be noted that HPV infections observed at the end of follow-up could be related to persistent infection, but also to new or recurrent infections. It was not possible to differentiate between them and this measurement bias cannot be addressed.

In 1997, Bollen et al. showed that HPV prevalence decreased from 98% to 31% when comparing samples before and after cervical treatment [15]. In that study, the five women living with HIV (not under antiretroviral therapy) all had positive HPV carriage after treatment. That study also indicated that detection of HPV infection after treatment was associated with incomplete conization. In our study, by the end of the follow-up, women with cervical abnormalities had a higher rate of HR-HPV infection, meaning that cervical lesions could be a proper medium for the persistence of HR-HPV infection. Also, in our study, all women were receiving effective antiretroviral therapy, and 97% had a CD4 level above 250 cells/mm³. This may have mitigated the mechanical effect of cervical treatment.

In a study in Nicaragua including 67 HIV-naïve women receiving LEEP for CIN2+ cervical lesions, 93% had at least one HR-HPV infection with genotypes 16, 31, 52, or 58 before treatment. Two years after treatment, the prevalence of HR-HPV infection was 8.4%, while it was 7.5% for CIN2+ cervical lesions [12]. In our study, the higher prevalence of HR-HPV carriage after treatment (27%) may be related to HIV infection, known to be a driver of incident HPV infections [1–3].

In a large study in China among HR-HPV infected women (HIV uninfected) without high-grade cervical lesions, viral clearance was estimated at 54.3% over a mean duration of 14.5 months, with higher clearance among women aged under 50 [13]. Evidence for age as a predictor of HPV clearance has been provided elsewhere [15]. This relationship reflects the natural course of HPV infection when ageing. In our study, probably due to lack of power, such association could not be evidenced.

Regardless of treatment, Travassos et al. showed a 46% clearance HR-HPV carriage after one year, among 62 women living with HIV and receiving antiretroviral therapy for more than a year [16]. This is consistent with the decrease we observed among women who received treatment (60%) and those who did not (30%). In our analysis, HPV 52 was the most frequent HR-HPV genotype evidenced both at the time of treatment indication and at the end of the follow-up. This high frequency was also evidenced among perinatally infected women in Thailand and Vietnam [7], emphasizing the need to scale-up nonavalent HPV vaccination programs in South-East Asia, at least among women living with HIV. The ongoing development of new L2-based vaccines may be promising for overcoming the diversity of oncogenic and potentially oncogenic HPV genotypes [17].

Our analysis is limited by a lack of statistical power, estimated post hoc at 16%. However, we did find that HR-HPV carriage decreased significantly over time among women who had received treatment.

5. Conclusions

Treatment of cervical lesions in women living with HIV under antiretroviral therapy may reduce the carriage of HR-HPV infections. Larger controlled cohort studies or meta-analyses are needed to assess which factors might be linked to a reduction of HR-HPV carriage among

women with cervical lesions. In this population, nonavalent or new L2-based vaccination programs could be considered for preventing HPV-induced cervical cancer.

Author Contributions: Conceptualization and methodology: S.L.C. and T.D. (Tristan Delory); implementation and investigation: C.P., P.S., W.M., P.L., S.B. and K.S.; biological analysis: N.N.-G.-H. and S.R.; statistical analysis: T.D. (Timothée Dub); writing—original draft preparation: T.D. (Timothée Dub); writing—initial review and editing: S.L.C. and T.D. (Tristan Delory); writing—final review, editing and agreement: All authors. All authors have read and agreed to the published version of the manuscript.

Funding: The study was supported by the Institut National du Cancer (INCa, Grant N° 2011-169); the Fondation de France (Grant N°00023970); the Global Fund to Fight AIDS, Tuberculosis and Malaria, Thailand (PR-A-N-008); the Ministry of Public Health, Thailand; and the Institut de Recherche pour le Développement (IRD), France; the Institut National d'Études Démographiques, France; and the Thailand International Development Cooperation Agency (TICA).

Informed Consent Statement: Informed consent was obtained from all subjects involved in the study.

Data Availability Statement: The data presented in this study are available on request from the corresponding author. The data cannot be made publicly available due to patients' confidentiality.

Acknowledgments: All the women who participated in the study; The PHPT team: Administrative support: Nitima Chaiboonruang, Pongsak Pirom, and Sakawrut Jitharidkul; Tracking & Regulatory: Kanchana Than-in-at, and Rukchanok Peongjakta; Safety monitoring: Suwalai Chalermpanmetagul; Sites monitoring: Pra-ornsuda Sukrakanchana, Cheeraya Kanabkaew, Boonyavee Ratchanee, Jirad-sadaporn Khanmali, Namthip Kruenua, Patcharaporn Krueuangkam, Renoo Kaewsai, Ruethai Wongchai, Subenya Jinasa, Tiwacha Thimakam, Waropat Pongchaisit, Warunee Khamjakkaew, Sirirat Thammajitsagul, Praewpan Roysuwan, and Boontila Insuyah. Data management: Sanuphong Chailoet, and Kriengkrai Suebmongkolchai; Laboratory: Nantawan Wangsaeng, Sirithorn Surajinda and Ampika Kaewbundit; IT: Luc Decker; The DSMB members: Heather Cubie, Silvia Franceschi, Marleen Praet, and Isabelle Heard; Christopher Leichtnam for his editorial expertise.

Conflicts of Interest: The authors declare no conflict of interest.

Appendix A

Table A1. Prevalence of cervical lesions, biopsy results, and Pap smear results before treatment and by the end of the follow-up among women who received treatment.

	Before Treatment Indication	Last HPV Sample	p Value
Presence of a cervical lesion	30 (100%)	2 (7%)	<0.001
Biopsy results			
- Squamous metaplasia	0 (0%)	2 (33%)	0.026
- CIN 1	1 (3%)	4 (67%)	
- CIN 2	10 (33%)	0 (0%)	
- CIN 3	19 (63%)	0 (0%)	
Pap smear results			

Table A1. Cont.

	Before Treatment Indication	Last HPV Sample	<i>p</i> Value
- Normal	10 (33%)	27 (90%)	0.001
- ASC-US	7 (23%)	1 (3%)	
- ASC-H	0 (0%)	0 (0%)	
- LSIL	3 (10%)	0 (0%)	
- HSIL	9 (30%)	2 (7%)	
- Squamous cell carcinoma	0 (0%)	0 (0%)	
- Atypical glandular cells	0 (0%)	0 (0%)	
- Adenocarcinoma <i>in situ</i>	0 (0%)	0 (0%)	
- Unsatisfactory	1 (3%)	0 (0%)	

Table A2. Prevalence of cervical lesions, biopsy results, and pap smear results at the time of treatment indication and by the end of the follow-up among women who did not receive treatment.

	Before Treatment Indication	Last HPV Sample	<i>p</i> Value
Presence of a cervical lesion	16 (100)	2 (13%)	<0.001
Biopsy results			
- Normal biopsy	2 (18)	1 (20%)	0.454
- Squamous metaplasia	1 (9)		
- CIN 1	5 (45)	3 (60%)	
- CIN 2	2 (18)		
- CIN 3	1 (9)	1 (20%)	

Table A2. Cont.

Before Treatment Indication		Last HPV Sample	<i>p</i> Value
Pap smear results			
-	Normal	2 (13)	12 (75%)
-	ASC-US		
-	ASC-H	1 (6%)	
-	LSIL	1 (6)	2 (13%)
-	HSIL	5 (31)	1 (6%)
-	Squamous cell carcinoma	1 (6)	0 (0%)
-	Atypical glandular cells	6 (38)	0 (0%)
-	Adenocarcinoma in situ	1 (6)	0 (0%)
-	Unsatisfactory		

<0.001

Table A3. Factors associated to the presence of an high-risk HPV infection by the end of the follow up among women presenting with a cervical lesion. Bivariate analyses adjusted on treatment.

	HR-HPV- (<i>n</i> = 32)	HR HPV+ (<i>n</i> = 14)	Total	aOR (IC95%) *	<i>p</i> -Value
Characteristics at treatment indication					
Age	40.4 (33.4–44.4)	37.4 (34.6–40.7)	39.4 (33.6–42.9)	1.0 (0.9–1.1)	0.801
CD4 cell count					
- >250 cells/mm ³	32 (100)	12 (86)	44 (96)	1	0.20
- <250 cells/mm ³	0	2 (14)	2 (14)	5.3 (0.4–+∞)	

Table A3. Cont.

	HR-HPV– (<i>n</i> = 32)	HR HPV+ (<i>n</i> = 14)	Total	aOR (IC95%] *	<i>p</i> -Value
HIV load					
-	<1.7 log10 copies/mL	29 (91)	13 (93)	42 (93)	1
					1.00
-	>1.7 log10 copies/mL	3 (9)	1 (7)	1 (7)	
HIV characteristics					
CD4 cell count at Nadir					
-	>250 cells/mm ³	4 (13)	0	4 (9)	1
					0.47
-	<250 cells/mm ³	28 (83)	14 (100)	42 (96)	
Time on ART at treatment indication					
	2.6 (1.3–4.0)	3.1 (1.4–6.6)	2.9 (1.4–4.6)	1.1 (0.9–1.3)	0.545
Last cytology					
-	Normal	32 (100)	7 (50)	39 (85)	1
					<0.001
-	ASC-US+	0	7 (50)	7 (15)	

* aOR: Odds ratio adjusted for treatment.

References

- Kantathavorn, N.; Mahidol, C.; Sritana, N.; Sricharunrat, T.; Phoolcharoen, N.; Auewarakul, C.; Teerayathanakul, N.; Taepisitpong, C.; Saeloo, S.; Sornsamtrang, G.; et al. Genotypic distribution of human papillomavirus (HPV) and cervical cytology findings in 5906 Thai women undergoing cervical cancer screening programs. *Infect. Agents Cancer* **2015**, *10*, 7. [\[CrossRef\]](#) [\[PubMed\]](#)
- Marks, M.A.; Gupta, S.; Liaw, K.-L.; Tadesse, A.; Kim, E.; Phongnarisorn, C.; Wootipoom, V.; Yuenyao, P.; Vipupinyo, C.; Rugpao, S.; et al. Prevalence and correlates of HPV among women attending family-planning clinics in Thailand. *BMC Infect. Dis.* **2015**, *15*, 159. [\[CrossRef\]](#) [\[PubMed\]](#)
- Phoolcharoen, N.; Kantathavorn, N.; Sricharunrat, T.; Saeloo, S.; Krongthong, W. A population-based study of cervical cytology findings and human papillomavirus infection in a suburban area of Thailand. *Gynecol. Oncol. Rep.* **2017**, *21*, 73–77. [\[CrossRef\]](#) [\[PubMed\]](#)
- Delory, T.; Ngo-Giang-Huong, N.; Rangdaeng, S.; Chotivanich, N.; Limtrakul, A.; Putiyanun, C.; Suriyachai, P.; Matanasarawut, W.; Jarupanich, T.; Liampongsabuddhi, P.; et al. Human Papillomavirus infection and cervical lesions in HIV infected women on antiretroviral treatment in Thailand. *J. Infect.* **2017**, *74*, 501–511. [\[CrossRef\]](#) [\[PubMed\]](#)
- Bollen, L.J.M.; Chuachoowong, R.; Kilmarx, P.; Mock, P.A.; Culnane, M.; Skunodom, N.; Chaowanachan, T.; Jetswang, B.; Neeyapun, K.; Asavapiriyant, S.; et al. Human Papillomavirus (HPV) Detection among Human Immunodeficiency Virus–Infected Pregnant Thai Women: Implications for Future HPV Immunization. *Sex. Transm. Dis.* **2006**, *33*, 259–264. [\[CrossRef\]](#) [\[PubMed\]](#)
- Ramautarsing, R.A.; Phanuphak, N.; Chaithongwongwatthana, S.; Wit, F.W.; Teeratakulpisarn, N.; Pankam, T.; Wijgert, J.H. Cervical and anal HPV infection: Cytological and histological abnormalities in HIV-infected women in Thailand. *J. Virus Erad.* **2015**, *1*, 96–102. [\[CrossRef\]](#)
- Sohn, A.H.; Kerr, S.; Hansudewechakul, R.; Gatechompol, S.; Choekphaibulkit, K.; Dang, H.L.D.; Tran, D.N.H.; Achalapong, J.; Teeratakulpisarn, N.; Chalermchokcharoenkit, A.; et al. Risk Factors for Human Papillomavirus Infection and Abnormal Cervical Cytology Among Perinatally Human Immunodeficiency Virus-Infected and Uninfected Asian Youth. *Clin. Infect. Dis.* **2018**, *67*, 606–613. [\[CrossRef\]](#) [\[PubMed\]](#)
- Verrier, F.; Le Coeur, S.; Delory, T. Cervical Human Papillomavirus Infection (HPV) and High Oncogenic Risk Genotypes among Women Living with HIV in Asia: A Meta-Analysis. *J. Clin. Med.* **2021**, *10*, 1911. [\[CrossRef\]](#) [\[PubMed\]](#)

9. Petry, K.U.; Horn, J.; Luyten, A.; Mikolajczyk, R.T. Punch biopsies shorten time to clearance of high-risk human papillomavirus infections of the uterine cervix. *BMC Cancer* **2018**, *18*, 318. [[CrossRef](#)] [[PubMed](#)]
10. Bollen, L.J.M.; Tjong-A-Hung, S.P.; Van Der Velden, J.; Mol, B.W.; Boer, K.; Kate, F.J.W.T.; Bleker, O.P.; Ter Schegget, J. Clearance of Cervical Human Papillomavirus Infection by Treatment for Cervical Dysplasia. *Sex. Transm. Dis.* **1997**, *24*, 456–460. [[CrossRef](#)] [[PubMed](#)]
11. Adebamowo, S.N.; Famooto, A.; Dareng, E.; Olawande, O.; Olaniyan, O.; Offiong, R.; Adebamowo, C.; ACCME Research Group as part of the H3Africa Consortium. Clearance of Type-Specific, Low-Risk, and High-Risk Cervical Human Papillomavirus Infections in HIV-Negative and HIV-Positive Women. *J. Glob. Oncol.* **2018**, *4*, 1–12. [[CrossRef](#)] [[PubMed](#)]
12. Aerssens, A.; Claeys, P.; Garcia, A.; Sturtewagen, Y.; Velasquez, R.; Broeck, D.V.; Vansteelandt, S.; Temmerman, M.; Cuvelier, C.A. Natural history and clearance of HPV after treatment of precancerous cervical lesions. *Histopathology* **2008**, *52*, 381–386. [[CrossRef](#)] [[PubMed](#)]
13. Guo, Y.-L.; You, K.; Qiao, J.; Zhao, Y.-M.; Liu, C.-R.; Geng, L. Natural history of infections with high-risk HPV in Chinese women with abnormal cervical cytology findings at baseline. *Int. J. Gynecol. Obstet.* **2010**, *110*, 137–140. [[CrossRef](#)] [[PubMed](#)]
14. Fregonese, F.; Collins, I.J.; Jourdain, G.; LeCoeur, S.; Cressey, T.R.; Ngo-Giang-Huong, N.; Banchongkit, S.; Chutanunta, A.; Techapornroong, M.; Lallemant, M. Predictors of 5-Year Mortality in HIV-Infected Adults Starting Highly Active Antiretroviral Therapy in Thailand. *JAIDS J. Acquir. Immune Defic. Syndr.* **2012**, *60*, 91–98. [[CrossRef](#)] [[PubMed](#)]
15. Costa, S.; De Simone, P.; Venturoli, S.; Cricca, M.; Zerbini, M.L.; Musiani, M.; Terzano, P.; Santini, D.; Cristiani, P.; Syrjänen, S.; et al. Factors predicting human papillomavirus clearance in cervical intraepithelial neoplasia lesions treated by conization. *Gynecol. Oncol.* **2003**, *90*, 358–365. [[CrossRef](#)]
16. Travassos, A.G.; Netto, E.; Xavier-Souza, E.; Nóbrega, I.; Adami, K.; Timbó, M.; Abbehusen, K.; Fernandes, S.; Duran, C.; Hagiuhara, T.; et al. Predictors of HPV incidence and clearance in a cohort of Brazilian HIV-infected women. *PLoS ONE* **2017**, *12*, e0185423. [[CrossRef](#)] [[PubMed](#)]
17. Huber, B.; Wang, J.; Roden, R.; Kirnbauer, R. RG1-VLP and Other L2-Based, Broad-Spectrum HPV Vaccine Candidates. *J. Clin. Med.* **2021**, *10*, 1044. [[CrossRef](#)] [[PubMed](#)]