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Vaginal Neoplasia Induced by an Unusual Papillomavirus Subtype in a Woman with Inherited Chromosomally Integrated Human Herpesvirus Type 6A

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Established Facts

- Human papillomaviruses (HPV) constitute a very heterogeneous family associated with cellular-proliferation pathologies.
- Systematic screening is recommended to diagnose cervical cancer. Cellular atypias were clearly associated with HPV infections of epithelial cells.
- Human herpesvirus type 6 (HHV6) could be integrated in cell genome and transmitted from parents to children.

Novel Insights

- HPV-4 was not known to induce severe neoplasia, especially in the genital tract of women.
- Inherited Chromosomally Integrated HHV-6A (iciHHV-6A) associated to low-risk HPV could be linked to tumoral development.

Keywords

Human papillomaviruses · Inherited chromosomally integrated HHV-6A · Low-grade SIL · High-grade SIL · Vaginal intra-epithelial neoplasia · Genital cancer

Abstract

We describe here a case of high-grade vaginal squamous lesion in a 54-year-old woman with a papillomaviruses (HPV) genital infection that developed from a cervical low-grade

squamous intraepithelial lesion (SIL) to a high-grade SIL (H-SIL) on cytological examination. A colposcopy exam led to the detection of suspect vaginal lesions with granulomatous infiltrations, which were classified as a Vaginal Intra-Epithelial Neoplasia grade 2 after pathologists' analyses. After a laser vaginal surgery and a loop excision of the transformation zone, the analyses of the anatomical pieces using a near-complete HPV screening panel revealed an HPV-4 infection that was not detected before in cervical smears. This HPV-infection is associated with a high human herpesvirus type 6A (HHV-6A) viral load in the same anatomical piece. The presence of an inherited chromosomally integrated HHV-6A (iciHHV-6A) was proved in this patient by real-time polymerase chain reaction on hair follicles and nail. This case suggests reconsidering both the benign nature of low-grade lesions in the female genital tract and the well-known "good" prognosis of low-risk HPV infection, especially when iciHHV-6A is diagnosed. This clinical course insists on the benefits of the multiplex panel use or global sequencing in order to optimize biological testing sensitivity, and so enhance clinical management of infection-induced neoplasia.

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Introduction

Newly developed molecular techniques have allowed discovering more than 170 human papillomaviruses (HPV) genotypes [1]. These viruses are implicated in cutaneous or mucosal lesions responsible for 4.8% of cancer worldwide [2]. HPV genotypes are divided into "high-risks" HPV responsible for HPV-induced cancers justifying a systematic screening and low-risk HPVs responsible for warts, papilloma, and low-grade squamous intraepithelial lesions (L-SIL) [1].

Inherited chromosomally integrated human herpesvirus type 6 (iciHHV-6) is rare and defined as when viral genome from HHV6 is integrated into the host germ line genome [3]. A positive polymerase chain reaction in nail and hair follicles allow us to distinguish infection by HHV6 and iciHHV-6 [3]. Spontaneous reactivation from iciHHV-6A was already observed in an immunosuppressed pediatric patient or after using valproate acid and carbamazepine [4].

We report here an atypical case of Vaginal Intra-epithelial Neoplasia (VaIN) grade 2 due to a low-risk HPV4 associated in a woman with an iciHHV-6A.

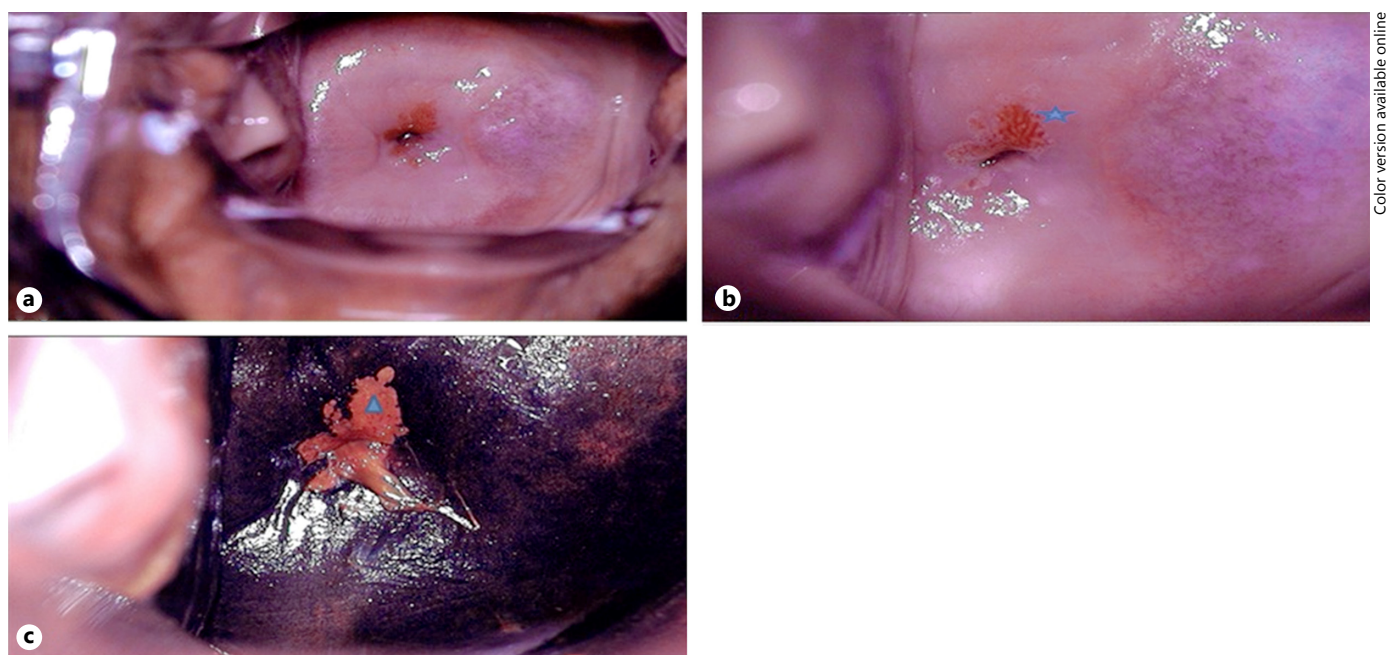


Fig. 1. Macroscopic aspect of the colposcopy. **a** Colposcopic aspect without preparation. **b** Colposcopic aspect after acetic acid application. **c** Aspect after lugol application. Star indicates the squamous junction area. Triangle indicates the iodopositive area.

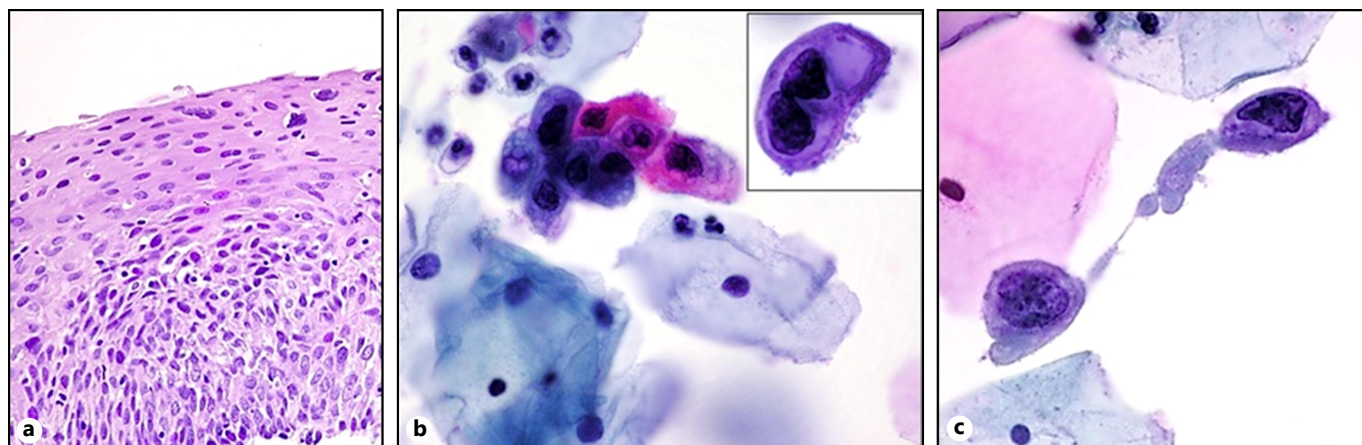


Fig. 2. Microscopic aspect of suspected neoplasia. **a** X400 VaIN2 aspect: chorionic inflammation associated with infiltration by lymphocytes and plasmocytes. The epithelial structure was modified on 1/3 to 1/2 of its length. **b** X1.000 Koilocytes aspect: small dichromatic basal cells with hyper chromatic nuclei. **c** X1.000 H-SIL aspect: small basal cells with irregular nuclei and cytoplasm.

Table 1. Patient's immunity exploration

a Cellular and humoral immunity exploration. Analyses were performed on April – May 2015 on Cell-Dyn(r) Sapphire system (Abbott, USA), FC500® (Beckman Coulter, Switzerland) and Hydrasys(r) (Sebia, France). No abnormalities were found on cellular quantification of leukocytes and lymphocytes and on quantification of immunoglobulins.

	Neutrophils, G/L	Lymphocytes, G/L	T4+, %	T8+, %	IgG, g/L	IgM, g/L	Phoresis
Results	3.61	2.01	52.36	15.47	8.29	0.78	Normal
Norm	1.5–8	1.5–6.5	31–63	12–37	7–16	0.4–2.8	–

G/L, giga per liter; T4+, TCD4+ cells; T8+, TCD8+ cells; IgG, immunoglobulin G; IgM, immunoglobulin M.

b Serological status of main viral infections. Analyses were performed using Liason® XL (DiaSorin, Italy).

	HSV 1–2	CMV	EBV	HHV-6	PVB19	Measles virus	Mumps virus	Rubella virus
IgG	+	+	+	+	+	+	+	+
IgM	–	–	–	–	–	–	–	NP

NP, non-performed.

* IgG are positive against VCA and EBNA for EBV serology.

Case Report

A 54-year-old woman under hormone replacement therapy, showed during systematic monitoring in 2002 cervical lesions classified as L-SIL. In 2008, an aggravation of the cytological aspect was found during monitoring justifying a colposcopy examination. An L-SIL at microscopic scale was found without abnormalities at macroscopic scale. No koilocytes, cellular reflects of the cytopathogenic effect caused by HPV infection, were observed during the pathologist's examination. In 2010, koilocytes were finally found.

In 2012, some lesions were classified as high-grade SIL, justifying exploration of the genital tract. During a complete vaginocopy, multiple suspect vaginal lesions and a cervix macroscopically normal were biopsied (Fig. 1), but no abnormalities were found on the hysteroscopy. Pathologists diagnosed a VaIN grade 2 on the histopathological aspect of these biopsies, confirmed by an augmentation of P16 protein expression and of the proliferation index (Ki-67) (Fig. 2a). Koilocytes were found on all samples (Fig. 2b, c). During all the clinical course (10 years) the search for infection by 35 HPV types (CLART2, Genomica, Madrid, Spain) remained

negative even on samples with koilocytes. Loop excision of the transformation zone and surgery were performed leading to the complete disappearance of infection and malignancy criteria. The patient is still closely monitored, but no sign of lesion has been observed so far.

All biopsies were retrospectively analyzed, by the International Agency for Research on Cancer that performed an extensive HPV screening panel (Luminex, TX, USA) ($n = 105$ genotypes). HPV-4 was found positive just above the detection threshold. Systematic *Herpesviridae* screening panel (Luminex, TX, USA) found a high but not quantifiable amount of HHV-6A DNA. A *Polyomaviridae* panel (Luminex, TX, USA) remained entirely negative.

Immunity was explored as well as the complete serological status for viral infections. No decrease of leukocytes or lymphocytes was found (Cell-Dyn[®] Sapphire system, Abbott, USA) and quantification for the CD4+ and CD8+ T cells were normal (FC500[®], Beckman Coulter, Swiss; Table 1a). Electrophoresis of immunoglobulins (Hydrasys(r), Sebia, France) was normal. The patient was seropositive for HHV-6 as for HSV-1, HSV-2, EBV, Measles, Mumps, rubella viruses and ParvovirusB19 (Liason[®] XL, Dia-Sorin, Italy) and seronegative for HIV (Table 1b).

HHV-6 quantification was performed on hair follicles and nail and blood samples and it showed a high HHV-6 viral load (6.73, 6.67, and 6.62 log₁₀copies/10⁶-cells, respectively), confirming the presence of iciHHV-6A.

Discussion

This atypical course of a low-grade lesion diagnosed during a systematic monitoring finally revealed a VaIN grade 2. This case is the first describing the association of an infection by a low-risk HPV and iciHHV-6A during the development of high-grade neoplasia in a female genital tract.

Although HPV-4 is known to be the sixth cause of benign cutaneous lesion or pre-cancer state, it is not known

to be associated with high-grade mucosal neoplasia, especially in immune patients [1, 5].

Explorations of the biopsies demonstrated a non-expected high load of HHV-6A, which may be responsible for neurological disease (tropism for oligodendrocytes) or immune system alteration (tropism for T-lymphocytes and NK cells) [6, 7].

In this case, the association of HPV-4 and iciHHV-6A in the cell could be responsible for a synergistic effect leading to severe cellular abnormalities. Previous studies suggested that iciHHV-6 could help the development of another viral infection by the modification of cellular receptor expression or pathways leading to possible predisposition to neoplasia [6]. Other studies, using a mice model, showed that HHV-6 could be an oncogenic virus by itself [8].

Without a specific treatment plan, patients with iciHHV-6 need to be intensively surveyed to optimize clinical management of suspect lesions at an early time of the disease before it progresses into the cancer state.

Disclosure Statement

Authors have no conflicts of interest to disclose.

Ethics Statement

For the purpose of this study, patient confidentiality was strictly maintained. This study was approved by the Ethical Committee of Hospices Civils de Lyon. Patient's approval has been searched, obtained, and then transmitted to the editorial board of the journal.

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