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Do people living with HIV face more secondary cancers than general population: from the French CANCERVIH network

Les personnes vivant avec le VIH sont-elles plus confrontées à des seconds cancers que la population générale : travail issu du réseau CANCERVIH

Marianne Veyri¹, Armelle Lavolé², Sylvain Choquet³, Dominique Costagliola⁴, Caroline Solas⁵, Christine Katlama⁶, Isabelle Poizot-Martin⁷, Jean-Philippe Spano⁸

¹AP-HP, Hôpitaux Universitaires Pitié-Salpêtrière – Charles Foix, Oncologie médicale, Sorbonne Université, INSERM, Institut Pierre Louis d'Epidémiologie et de Santé Publique, équipe Theravir, 47-83 boulevard de l'Hôpital 75013 Paris, France ²Sorbonne Université, GRC n°4, Theranoscan, AP-HP, Service de Pneumologie, Hôpital Tenon, 4 rue de la Chine 75020 Paris, France ³AP-HP, Hôpitaux Universitaires Pitié-Salpêtrière – Charles Foix, Hématologie Clinique, 47-83 boulevard de l'Hôpital 75013 Paris, France ⁴Sorbonne Université, INSERM, IPLESP (Institut Pierre Louis d'Epidémiologie et de Santé Publique), 47-83 boulevard de l'Hôpital 75013 Paris, France ⁵Aix Marseille Univ, APHM, INSERM 1207, IRD 190, UVE, Hôpital La Timone, Laboratoire de Pharmacocinétique et Toxicologie, 264 rue Saint-Pierre 13005 Marseille, France ⁶Sorbonne Université, INSERM, Institut Pierre Louis d'Epidémiologie et de Santé Publique, équipe Theravir, AP-HP, Hôpitaux Universitaires Pitié Salpêtrière - Charles Foix, Maladies Infectieuses et Tropicales, 47-83 boulevard de l'Hôpital 75013 Paris, France ⁷Aix Marseille Univ, APHM, INSERM, IRD, SESSTIM, Sciences Economiques & Sociales de la Santé & Traitement de l'Information Médicale, Hôpital Sainte-Marguerite, Service d'Immuno-hématologie clinique, 270 boulevard de Sainte-Marguerite 13009 Marseille, France ⁸Sorbonne Université, INSERM, Institut Pierre Louis d'Epidémiologie et de Santé Publique, équipe Theravir, AP-HP, Hôpitaux Universitaires Pitié-Salpêtrière - Charles Foix, Oncologie médicale, 47-83 boulevard de l'Hôpital 75013 Paris, France

<u>Corresponding author:</u> Marianne Veyri, Medical Oncology department, Pitié-Salpêtrière Hospital, 47-83 boulevard de l'Hôpital, 75013 Paris, +33 (0)1 42 16 00 26 <u>marianne.veyri@aphp.fr</u>

ABSTRACT

Introduction: People living with HIV (PLWHIV) are at a higher risk of cancer compared to the general population. With improved cancer treatments and the increased life expectancy of PLWHIV, the incidence of second cancers is also expected to increase.

Methods: We reviewed the cases of PLWHIV with cancer that have been presented to the CANCERVIH national multidisciplinary board since 2014. We included all cases with a history of cancer, and studied the incidence and types of second cancers.

Results: 719 cases were reviewed, out of which 94 (13%) had a history of at least one cancer. For the first primary cancers, 46 (49%) were AIDS-defining cancers (ADCs) and 48 (51%) were non-AIDS-defining cancers (NADCs). Kaposi sarcoma (33%) and NHL (15%) occurred most frequently as first cancers.

Among the first cancers that were ADCs, 15% of the second cancers were NHL, 11% anal canal cancers, 9% bladder and 9% Hodgkin lymphomas. Among the first cancers that were NADCs, 38% of the second cancers were lung cancers, 8% bladder, 8% head and neck and 8% NHL.

Discussion: With the aging of PLWHIV, the incidence of second and subsequent cancers is expected to increase in this population. Immuno-virological control should be maintained. Increased surveillance, early prevention and screening programs should be offered to all PLWHIV, including those with an undetectable HIV viral load and/or immune restoration.

RESUME

Introduction : Les personnes vivant avec le VIH (PVVIH) ont un risque plus élevé de cancer que la population générale. Avec l'amélioration des traitements carcinologiques et l'augmentation de l'espérance de vie des PVVIH, l'incidence des seconds cancers devrait augmenter.

Méthodes : Nous avons examiné les dossiers des PVVIH atteintes d'un cancer qui ont été présentés en RCP nationale CANCERVIH depuis 2014. Nous avons inclus tous les cas avec antécédents de cancer, et étudié l'incidence et les types de seconds cancers.

Résultats : 719 dossiers ont été examinés dont 94 (13%) avaient un antécédent d'au moins un cancer. Pour les premiers cancers primitifs, 46 (49 %) étaient des cancers classant-SIDA (ADCs) et 48 (51 %) des cancers non-classant-SIDA (NADCs). Le sarcome de Kaposi (33 %) et le LNH (15 %) étaient les premiers cancers les plus fréquents.

Parmi les premiers cancers qui étaient des ADCs, 15 % des seconds cancers étaient des LNH, 11 % des cancers du canal anal, 9 % de vessie et 9 % des lymphomes de Hodgkin. Parmi les premiers cancers qui étaient des NADCs, 38% des seconds cancers étaient des cancers du poumon, 8% de vessie, 8% de la tête et cou et 8% des LNH.

Discussion : Avec le vieillissement des PVVIH, l'incidence des seconds cancers (et ultérieurs) devrait augmenter dans cette population. Le contrôle immuno-virologique doit être maintenu. Surveillance accrue, prévention précoce et programmes de dépistage devraient être proposés à toutes les PVVIH, même avec une charge virale VIH indétectable et/ou une restauration immunitaire.

Keywords: AIDS-defining cancers, non-AIDS-defining cancers, HIV, second cancer **Mots clés :** cancers classant-SIDA, cancers non-classant-SIDA, VIH, second cancer

This study was the subject of a poster at ESMO 2019 (# 2036P).

INTRODUCTION

The life expectancy of people living with HIV (PLWHIV) has considerably improved since the advent of effective antiretroviral therapy (ART) in 1996, approaching that of the general population^{1,2}. As a consequence, causes of mortality in PLWHIV have shifted from AIDS to non-AIDS causes, making non-AIDS-related cancers the leading cause of non-AIDS deaths in PLWHIV^{1;3;4;5}. PLWHIV are at increased risk of cancer compared to the general population, as a result of chronic immunosuppression, concomitant infection with oncogenic viruses (human papillomavirus [HPV], hepatitis B [HBV] and C [HCV] viruses, Epstein-Barr Virus [EBV]), as well as prevalence of smoking and alcohol use in the HIV-positive population^{6;7;8;9.} Cancers in PLWHIV are typically divided into AIDS-defining cancers, which are known to occur at increased rates and mark the onset of AIDS (ADCs: Kaposi sarcoma, non-Hodgkin lymphoma [NHL] and cervical cancer) and non-AIDS-defining cancers (NADCs). The incidence of ADCs has declined since the introduction of ART, but the risk of NADCs has increased with the growth and aging of the HIV-positive population^{4;8;10}.

Due to improved outcomes of cancer patients, the population of cancer survivors is growing, with a 5-year cancer prevalence of 1.39 million in France alone (Globocan 2018). Studies on the incidence of second cancers have shown that cancer survivors are at an increased risk of developing a new cancer compared to the general population, due in part to late effects of cancer treatments, but also etiologic and environmental factors^{11;12}. Due to the longer survival of PLWHIV, the incidence of secondary cancers is also expected to increase in PLWHIV. Furthermore, PLWHIV have been shown to have worse survival from cancer compared to non-infected individuals^{13;14}. Improvements in cancer prevention, screening and treatment are therefore crucial in high-risk populations such as PLWHIV who have had cancer. However, data is limited on the risk of second cancer in PLWHIV. In a study that assessed the incidence of second cancers in PLWHIV using the San Francisco HIV/AIDS case registry and California Cancer Registry, PLWHIV were shown to have an increased risk for both first and second primary cancers, before and in the era of effective ART, in particular Kaposi sarcoma, NHL, Hodgkin's lymphoma, anal cancer, and liver cancer¹⁵. This study also

showed that while first and second primary ADC incidence declined, second primary NADC incidence increased over time. Many of the cancers in excess had viral causes, or were related to smoking or alcohol use.

We conducted a study of a national prospective database of PLWHIV with cancer to evaluate the incidence of secondary cancer in PLWHIV, analyze the types of cancer and sequence of occurrence.

METHODS

Study design and data collection

This was an observational study to evaluate the incidence of secondary cancers in PLWHIV in the CancerVIH database. CancerVIH is a national multidisciplinary network dedicated to HIV-infected patients with cancer¹⁶ accredited by the French National Cancer Institute, and that has obtained all the necessary regulatory approvals (French IRB [n°15-009], CCTIRS [n°16-391] and the Commission Nationale de l'Informatique et les Libertés [CNIL; n°916500]). CancerVIH established a database in May 2014 to prospectively include new cases of PLWHIV with cancer. Each case is discussed during the bi-monthly national multidisciplinary board, composed of oncologists, radiotherapists, hematologists, infectiologists, virologists, immunologists and pharmacologists, in web-conference. Anonymized medical files are presented by the patients' referring physicians (HIV specialist or oncologist). Cases are presented on a regular basis by 109 centres in France (and a few centres in Switzerland which do not have a similar referral network).

We reviewed all cases in the CancerVIH database between 6 May 2014 (database creation date) and 31 December 2019.

For this study, we examined the history of cancer indicated by the referring physicians. History of cancer that was neither metastasis nor locoregional relapse was considered as primary cancer. We included in the analysis patients with prior cancer at a different site, at the same site with a different histology or a concomitant cancer at a different site or with a different histology.

Patients with more than one previous malignant disease were included.

Demographic, epidemiological and immuno-virological characteristics were described at the time of the last cancer diagnosis.

RESULTS

Patient characteristics

In total, 719 cancer cases were reviewed. All patients were aged 18 or older and 97% were on ART. Among all these cases, 94 patients (13%) had a history of at least one previous cancer. Patient characteristics are summarized in Table 1.

Of those 94 patients, 80 (84%) were men and 14 (16%) were women. All were under ART. The median age at onset of first cancer was 51 years for all patients (51 years for men, 48 years for women). The median age at onset of second cancer was 56 years for all patients (58 years for men, 52,5 years for women). The median CD4 nadir was 80 cells/mm³. At the time of diagnosis of the last cancer, the median time spent on ART was 15 years. The median CD4 count was 400 cells/mm³ and all patients were with undetectable HIV viral load. Among the 22 patients (23%) who had more than 2 cancers: 16 patients (17%) had 3 cancers, 3 patients (5%) had 4 cancers, 2 patients (2%) had 5 cancers and 1 patient (1%) had 7 cancers. The median time from first cancer to second cancer was 8,5 years.

Cancer types and sequence of occurrence

For the first primary cancers, 46 were ADCs (49%) and 48 (51%) were NADCs (Figure 1). Kaposi sarcoma (33%) and non-Hodgkin lymphoma (15%) occurred most frequently as first cancers. The most frequent first primary non-AIDS-defining cancers were head and neck (7%), breast (5%), Hodgkin lymphoma (5%), and prostate cancers (5%).

Among the first cancers that were AIDS-defining cancers, 15% of the second cancers were NHL, 11% anal canal cancers, 9% bladder cancers and 9% Hodgkin lymphomas. Among the first cancers that were non-AIDS-defining cancers, 38% of the second cancers were lung cancers, 8% bladder, 8% head and neck and 8% NHL.

DISCUSSION

In this study, we evaluated the incidence of second cancers in a population of PLWHIV with cancer in a national database. Overall, 13% of patients had second or subsequent cancers, a higher proportion than that found in the study on HIV-positive San Francisco residents diagnosed with cancer between 1985 and 2013 (9%, 372/4144)¹⁵. It was also higher than the incidence of second cancer (7.3%) in the general population found in a study using data from the French network of cancer registries (FRANCIM) of nearly 290,000 patients diagnosed with cancer between 1989 and 2004¹⁷ and that in another study using surveillance data of over 2 million patients in the U.S. diagnosed with one of the ten most common cancers between 1992 and 2008 (8%)¹⁸. Nevertheless, the proportion was getting closer to that of second cancers in the general population of cancer survivors in the U.S. according to Surveillance, Epidemiology, and End Results (14-18%^{11;12;19}).

The types of first cancer in this study were almost equally divided into ADCs and NADCs (49% and 51%, respectively). Kaposi sarcoma and NHL occurred most frequently as first cancer (33% and 15%, respectively). For those who had Kaposi sarcoma, the most common secondary cancers were NHL (19%) and anal canal cancer (13%). In the San Francisco registry study, 50% of the HIV-positive patients diagnosed with cancer between 1990 and 2010 had a Kaposi sarcoma and 20% NHL as first cancer¹¹.

The most frequent first primary NADCs were head and neck cancers (7%), breast (5%), prostate (5%), Hodgkin lymphoma (5%), kidney (4%), anal canal (3%) and melanoma (3%). Many of the NADCs either as first or subsequent cancer we observed in this study could be due, at least in part, to viral causes, notably ano-genital and oropharyngeal cancers (HPV) and Hodgkin lymphoma (EBV). Lifestyle factors, such as smoking and alcohol use, may also put PLWHIV at risk of second cancers. Among the most common NADCs that could be related to smoking and alcohol use were lung, head and neck, liver and pancreas. However, factors other than smoking may explain the high incidence of lung cancer in the HIV-positive population²⁰⁻²¹. Biomarkers need to be identified for efficient prevention of the most common second cancers such as anal cancer²², and interventions should be pursued

to encourage modification of cancer-associated lifestyles (smoking and alcohol consumption).

Immunosuppression caused by HIV infection and implicated in the development of certain cancers is increased by low CD4 nadir and prolonged duress without effective ARV treatment. ARV treatment should be started as early as possible and continued for life, even during cancer treatment, in order to maintain high CD4 counts and an undetectable viral load.

Our study has several limitations. This was a retrospective observational study with a relatively small sample size. Data on lifestyle habits (tobacco and/or alcohol consumption) and all previous treatments are not systematically collected and could not be studied. Nevertheless, patients were included in the database prospectively, and the database is not exhaustive knowing the fact that the incidence in France of cancer in PLWHIV is about 13‰¹. Referring physicians call indeed on the network for the most serious cases requiring an appeal decision. In addition, our study population was predominantly male (84%), thereby further limiting the sample size of female PLWHIV and their malignancies.

With the aging of PLWHIV, the incidence of second and subsequent cancers is expected to increase in this population, who are at risk for both ADCs and NADCs. Our findings strongly suggest the importance of effective screening and prevention of all types of cancers in PLWHIV. Immuno-virological control should be maintained in PLWHIV with cancer. Increased surveillance, early prevention including vaccination and screening programs should be offered to PLWHIV with previous cancer history, including those with an undetectable HIV viral load and/or immune restoration.

Table 1:	Patient ch	naracteristics

Patient characteristics	Number of patients
Sex	
Male	80 (84%)
Female	14 (16%)
Median age 1 st cancer (all genders) Male Female	50 years (IQR, 41,5-56) 51 years (IQR, 42-59) 48 years (IQR, 36,5-54)
Median age 2 nd cancer (all genders) Male Female	56 years (IQR, 51-64) 58 years (IQR, 51-64) 52,5 years (IQR, 46-64)
Median CD4 nadir	80 cells/mm ³ (IQR, 17-156,5)
Time exposure to ART	15 years (IQR, 9-23)
Median CD4	400 cells/ mm ³ (IQR, 222,5-651)
Median HIV VL	0 cp/mL (IQR, 0-0)
Year of discovery of HIV infection	
Before 1997	47 (50%)
1997-2000	19 (20%)
2001-2004	3 (3%)
2005-2009	12 (13%)
After 2009	12 (13%)
Undetermined	1 (1%)
Delay between HIV diagnosis	
and cancer diagnosis	
History of first cancer prior to HIV	4 (4%)
diagnosis	
Concomitant diagnosis of HIV	10 (11%)
and first cancer	
1 year after HIV diagnosis	6 (6%)
Between 2 and 5 years after	7 (7%)
Between 6 and 10 years after	10 (11%)
Between 11 and 20 years after	23 (24%)
More than 21 years later	11 (12%)
Data not filled in	23 (24%)
First primary cancers	
ADC	46 (49%)
NADC	48 (51%)
Second cancer with an ADC first	
primary cancer	
NHL	7 (15%)
Bladder cancer	5 (11%)
Canal anal cancer	5 (11%)
Hodgkin lymphoma	4 (9%)
Head and neck cancer	3 (7%)
Lung cancer	3 (7%)
Castleman disease	3 (7%)
Liver cancer	2 (4%)
Colorectal cancer	2 (4%)

Biliary cancer	2 (4%)
Other	10 (20%)
Second cancer with a NADC first	
primary cancer	
Lung cancer	18 (38%)
Bladder cancer	4 (8%)
Head and neck cancer	4 (8%)
NHL	4 (8%)
Kidney cancer	2 (4%)
Colorectal cancer	2 (4%)
Breast cancer	2 (4%)
Other	12 (25%)
Median time from 1st cancer to 2 nd cancer	8,5 years (IQR, 3-12)

Figure 1: description of the second cancers in relation to the first cancer

14 NHL	 Dedgkin lymphoma Hodgkin lymphoma - Dedge Anal canal cancer - Dedge Lung cancer Anal canal cancer Bladder cancer Myeloma Head and Neck cancer Lung cancer Liver cancer Liver cancer 	 Anal canal cancer Anal canal cancer Anal canal cancer Anal canal cancer - 3 Bowen disease NHL NHL NHL NHL NHL NHL NHL
	② Penis cancer	2 NHL
	② NHL - ③ Colorectal cancer	2 NHL
	② Prostate cancer - ③ NHL	② Castleman disease
	② Kidney cancer - ③ Hodgkin lymphoma	© Castleman disease
	② Vulva cancer - ③ Lung cancer	© Castleman disease
	② Melanoma - ③ Anal canal cancer - ④ Head and Neck cancer	© Liver cancer
		© Pancreas cancer
		© Lung cancer
		© Lung cancer
		© Colorectal cancer
		© Colorectal cancer
	31 Kaposi diseases	② Head and Neck cancer
		② Head and Neck cancer
		© Bladder cancer
		2 Bladder cancer
		© Bladder cancer
		© Biliary cancer
		© Biliary cancer
		② Anal margin cancer - ③ Lung cancer
		Cutaneous cancer - 3 Cutaneous cancer - 4 Lung cancer
		② Hodgkin lymphoma
	1 cervix cancer ^② Leiomyosarcoma	D Hodgkin lymphoma - D Anal canal cancer - Head and Neck cancer - Kaposi disease
		Urothelial cancer - 3 Cutaneous cancer - 9 Prostate cancer - 9 Melanoma - 9 Pancreas cancer - 9 Lung cancer

ADC (AIDS-defining cancer) as first cancer

7 Head and Neck	cancers	 © Esophagus cancer © Lung cancer © Lung cancer © Lung cancer © Bladder cancer © Head and Neck cance © Head and Neck cance 	er - ③ Lung cancer er - ③ Head and Neck cancer - ④ Head an	d Neck cancer - ① Head and Ne	5 Breast cancer	© Head and Neck cancer © NHL © Lung cancer © Lung cancer © Lung cancer
5 Prostate cance	 2 Colore 2 Colore 2 Leuker 2 Lung co 2 Bladde 	ectal cancer ectal cancer mia ancer er cancer		5 Hodgkir	n lymphomas	9 NHL 9 Lung cancer 9 Lung cancer 9 Lung cancer 9 Hodgkin lymphoma - ③ MDS
4 Kidney cancers	 C Kidney c Lung can Lung can C Stecho 	ancer cer Indroma - ③ Breast cance	3 Anal canal cancers	 Anal margin cancer Lung cancer Lung cancer Lung cancer 	2 Liver cancers ncers © Lung cancer © Lung cancer	 Uung cancer NHL - Lung cancer - Lung cancer
3 Melanomas	 Ø Melanoma Ø Breast canc Ø Bladder canc 	- ③ Bladder cancer er icer	2 Anal margin cancers	© Bowendisease © NHL 1 Colorectal	cancer © Kidney ca	ncer - ③ Colorectal cancer
2 Cutaneous can	cers © R	ectal cancer aposi disease - ③ Head a	nd Neck cancer 1 Gastric	Cancer © Bladder cancer	r - ③ Kaposi disease	
2 Eye cancers	② CML③ Meningiom	a - ③ Head and Neck can	1 Thyroid cancer	 Dead and Neck cancer 	1 Bowen	disease _{© CUP}
NADC (Non-AIDS-defining cancer) as first cancer						

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COMPLIANCE WITH ETHICAL STANDARDS

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Conflict of Interest: Veyri M, Lavolé A, Choquet S, Costagliola D, Solas C, Katlama C and Poizot-Marin I declare that they have no conflict of interest. Spano JP has no direct conflict of interest but received an honorarium as a consultant for

Roche, MSD and Biogaran and has received a speaker honorarium from MSD, Roche,

AstraZeneca, Leopharma, Mylan, Pfizer, BMS, Novartis, PFO, Myriads, Gilead and Lilly.

Ethical approval: This article does not contain any studies with human participants performed by any of the authors. This is a non-interventional study.

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