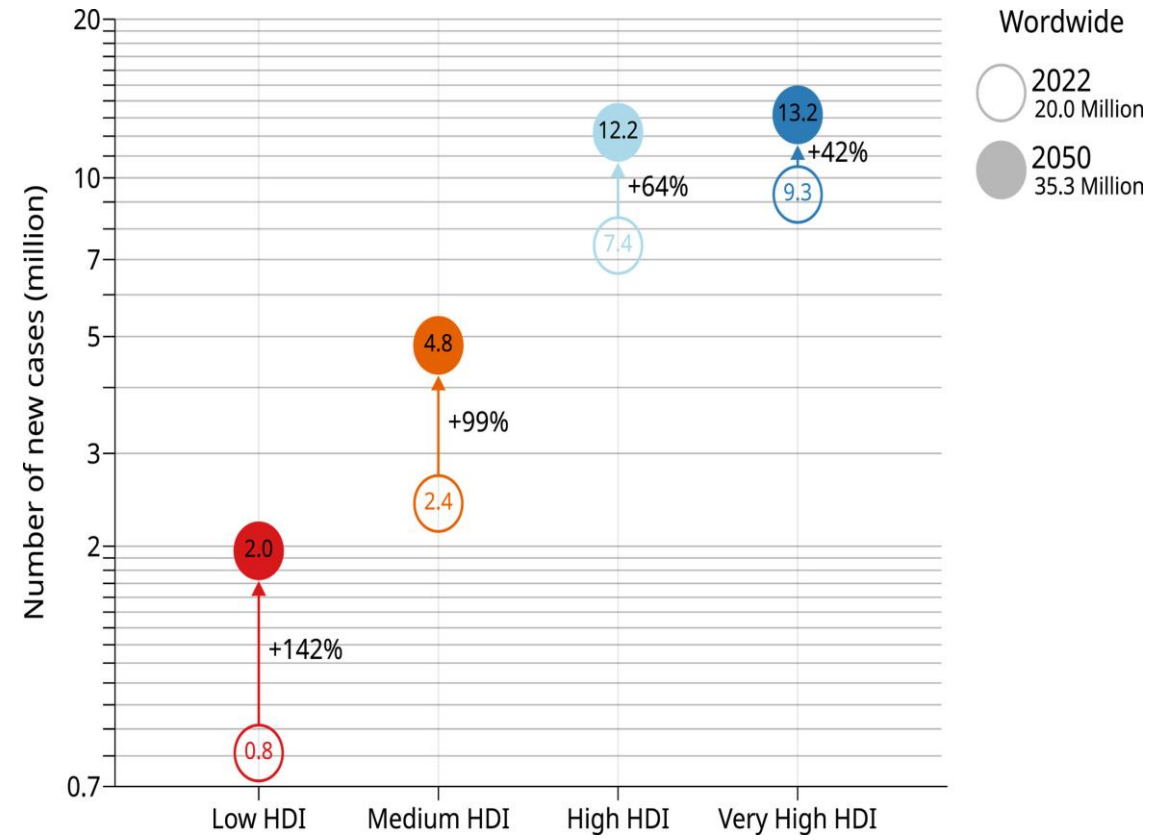
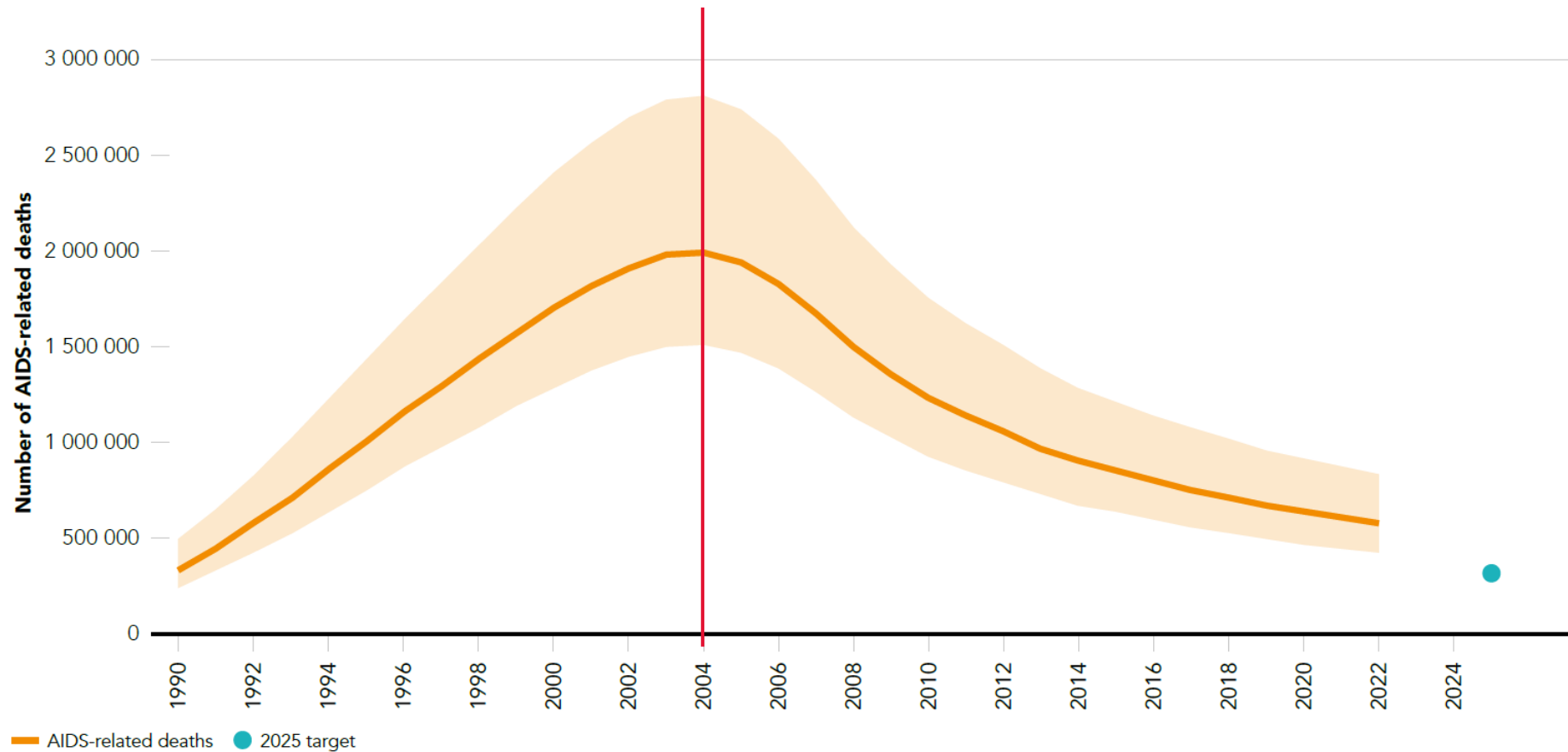


Source: UNAIDS epidemiological estimates, 2023 (<https://aidsinfo.unaids.org/>).

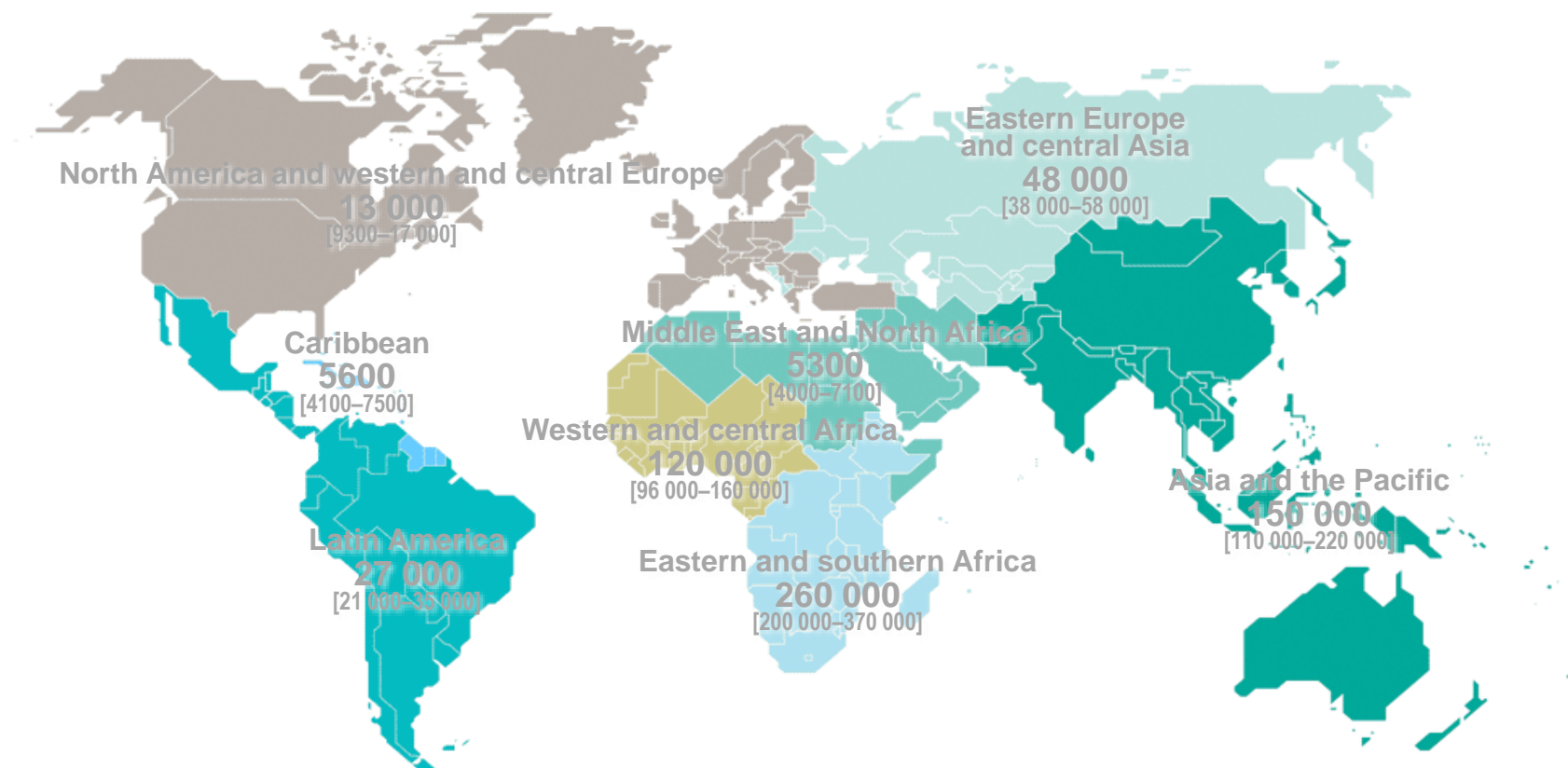


# Number of HIV-related deaths, global, 1990-2022



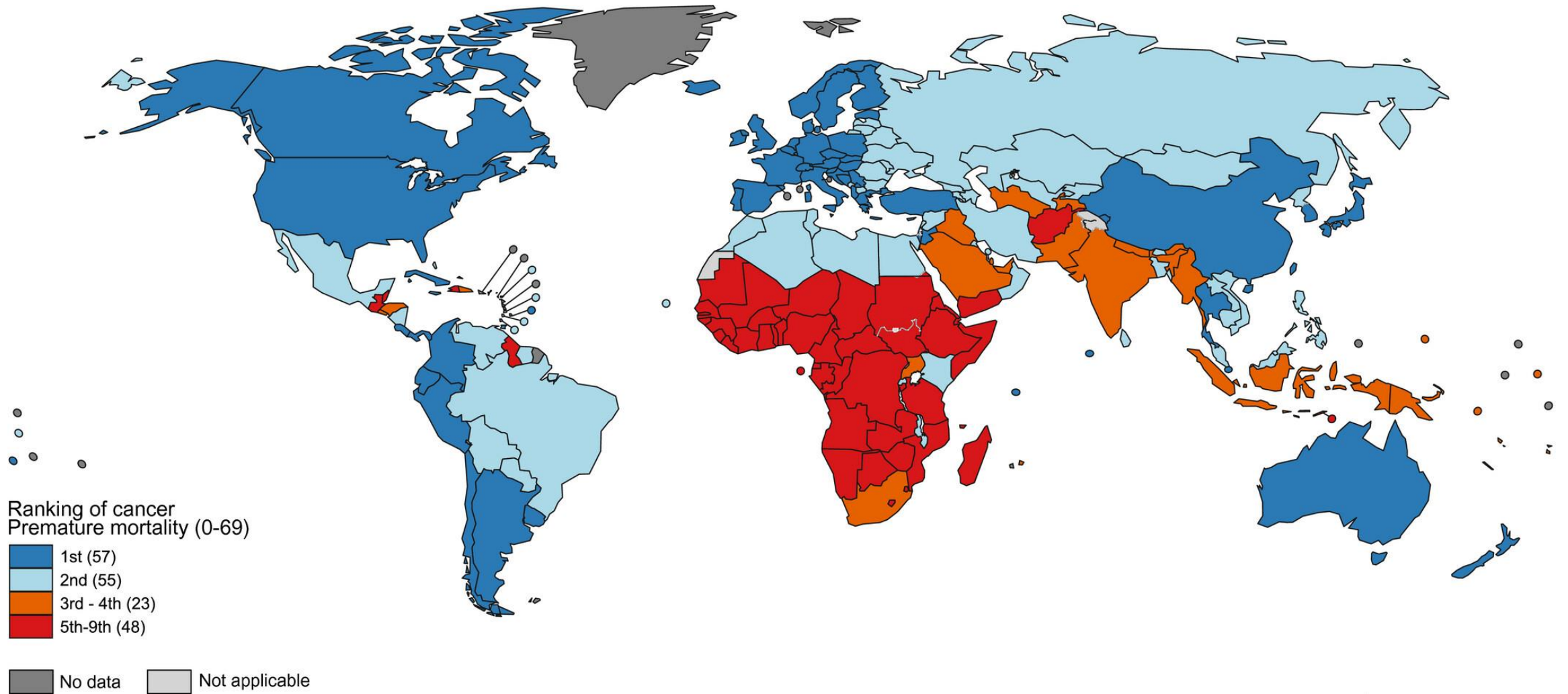
Source: UNAIDS epidemiological estimates, 2023 (<https://aidsinfo.unaids.org/>).

## Estimated adult and child deaths from AIDS | 2022



**Total: 630 000** [480 000–880 000]

# Global burden of cancer: today



# Global burden of cancer: today

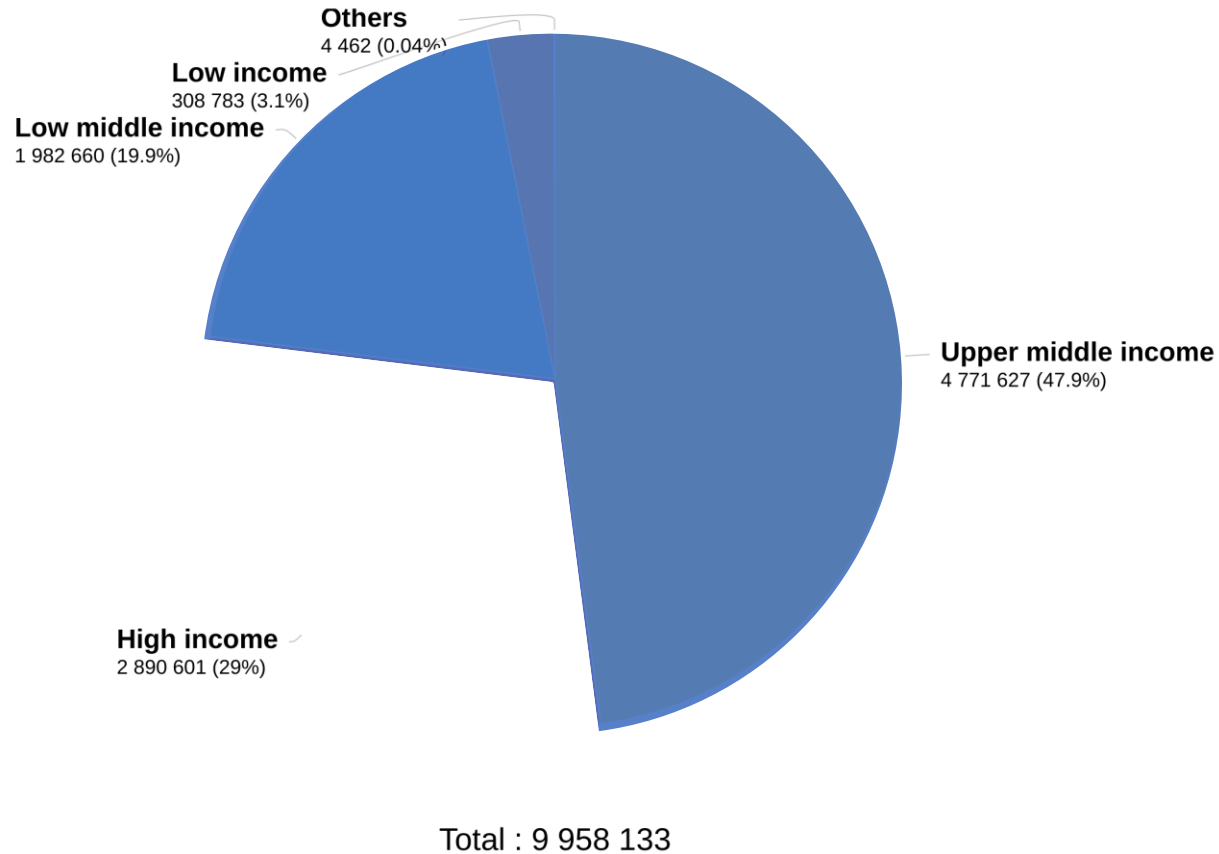
Estimated number of global cancer deaths in 2022

**9,736,779**

**10 million** cancer deaths in 2022

# Global burden of cancer: today

Estimated number of global cancer deaths in 2022



**10 million** cancer deaths in 2022

**70%** in low- and middle-income countries

# Beyond Barriers: Enhancing the Reach of HIV Service Delivery by **Addressing Cancer** **in PWH**

OR

***“We should put out the fire while it is still small.”*** – Kenyan proverb

**Thomas A. Odeny, MD, MPH, PhD**

Assistant Professor of Medicine  
Division of Oncology, Washington University in St. Louis

Senior Principal Clinical Research Scientist  
Kenya Medical Research Institute

▮ *“We should put out the fire while it is still small.” – Kenyan proverb*

# DISCLOSURES

1. Research funding from Gilead Sciences

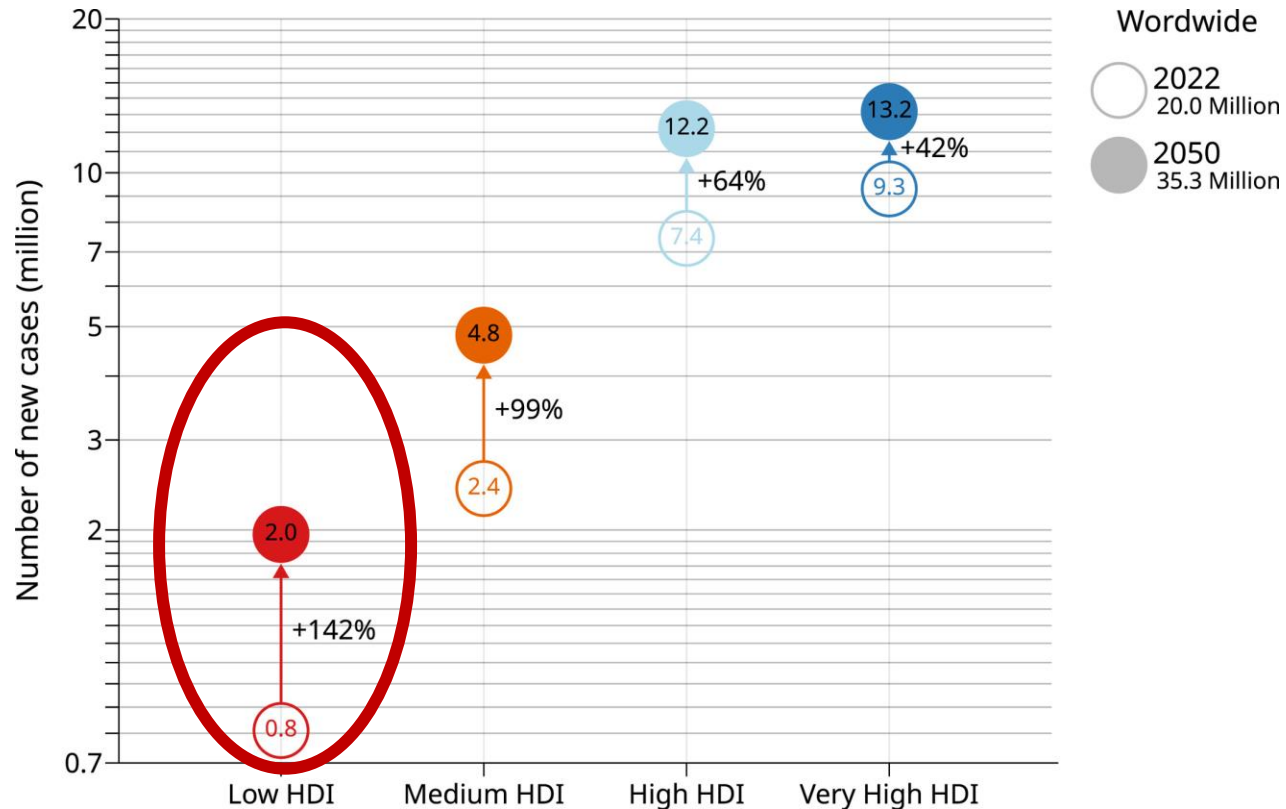


# OUTLINE

- 1. Global Burden of Cancer vs. HIV**
2. Cancer in People with HIV
3. Opportunities to Enhance the Reach of HIV Service Delivery by Addressing Cancer in PWH
4. Conclusions/Invitation

# Global burden of cancer 2022-2050

Estimated increase in new cancer cases 2022-2050

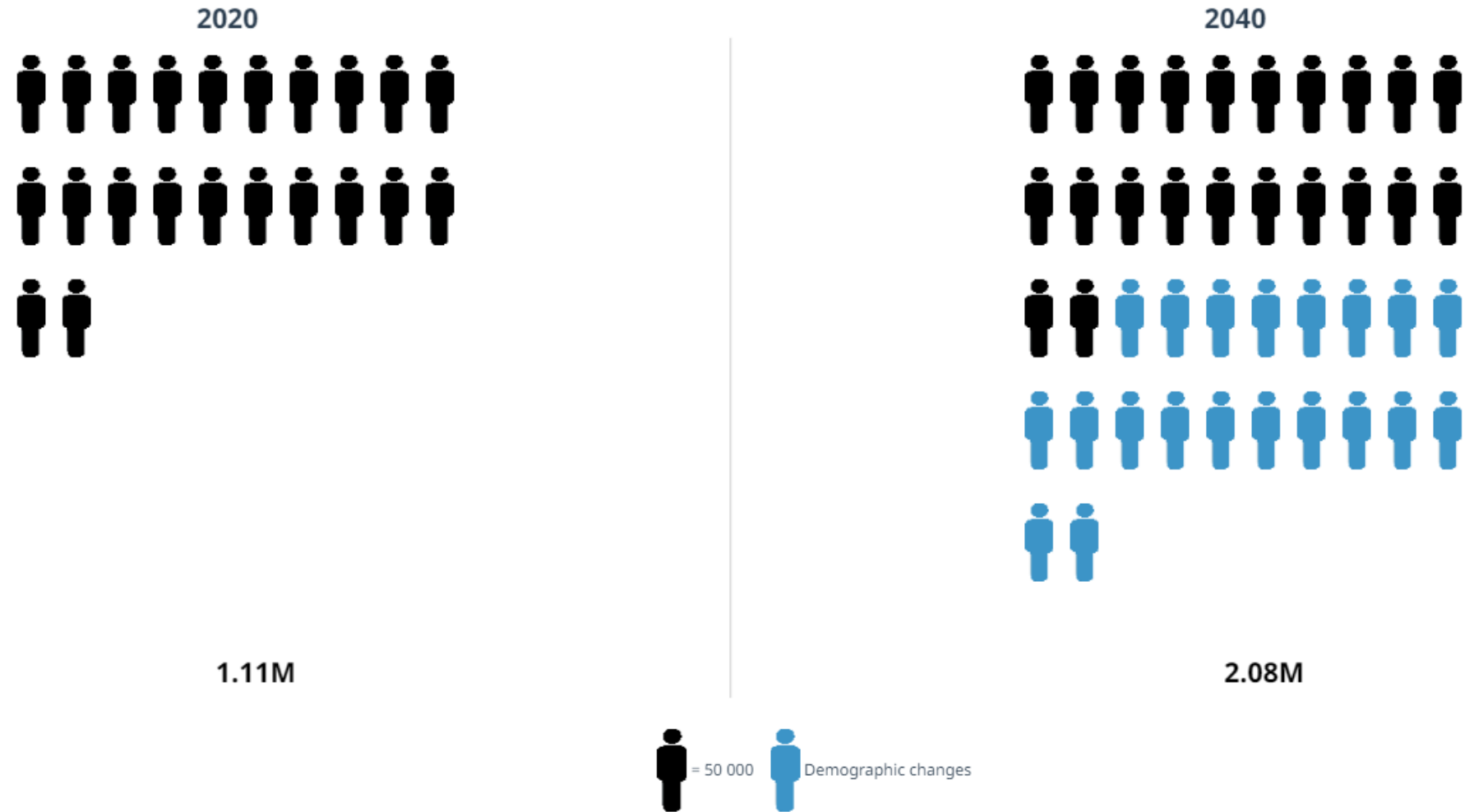


**35 million** new cancer cases by 2050

**142%** increase in new cancer cases in low-income countries

# Global burden of cancer: 2020-2040

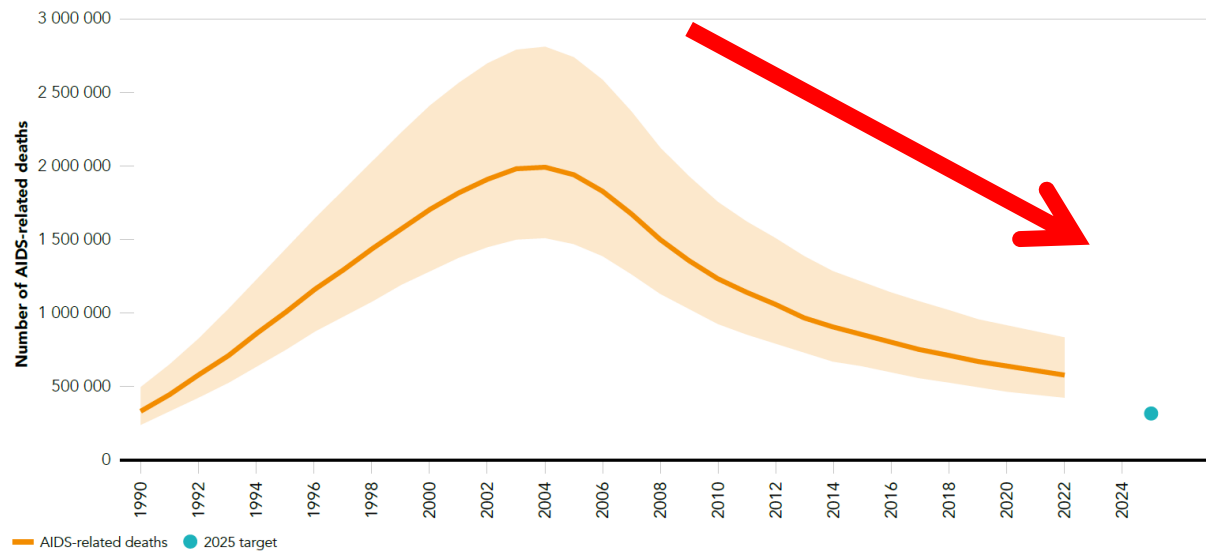
Estimated increase in new cancer cases 2020-2040, Africa



# Comparing Global Burden of HIV and Cancer

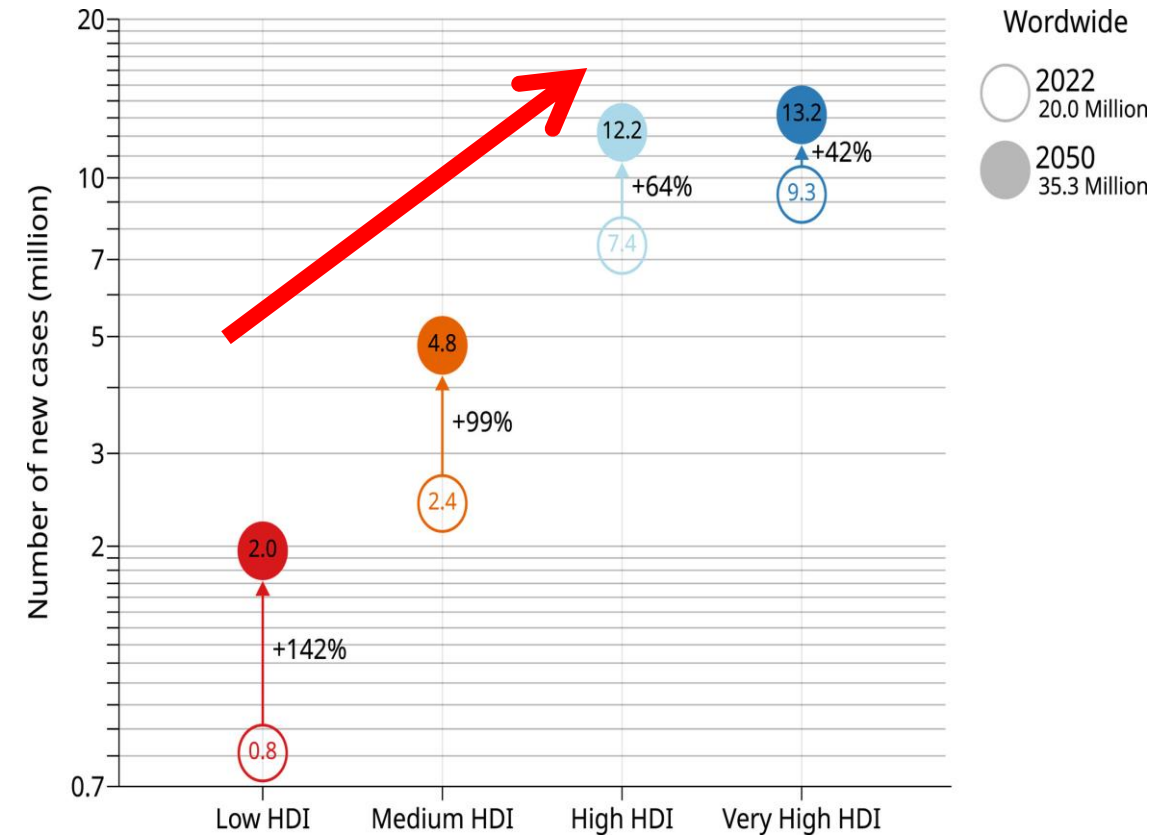
## HIV-related deaths

Figure 12.2 Number of AIDS-related deaths, global, 1990–2022, and 2025 target



Source: UNAIDS epidemiological estimates, 2023 (<https://aidsinfo.unaids.org/>).

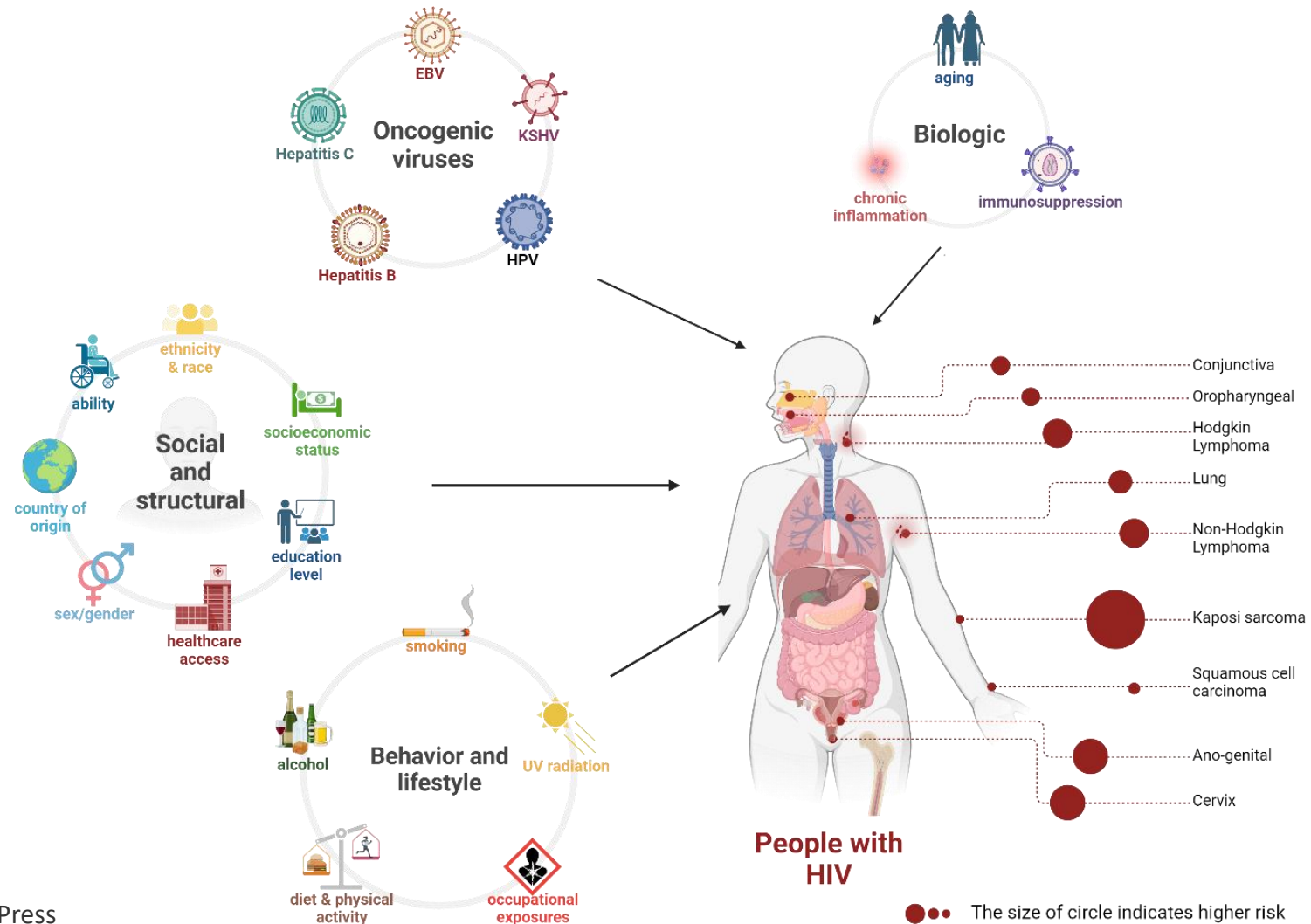
## Cancer-related deaths



# OUTLINE

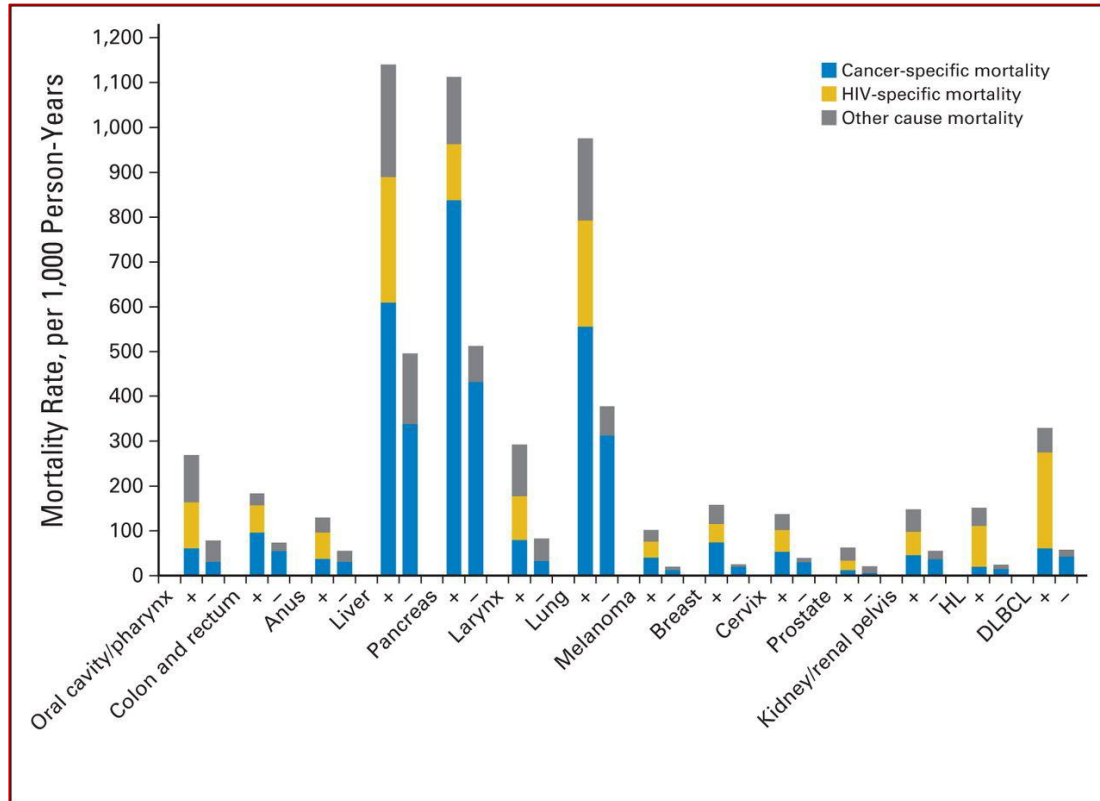
1. Global Burden of Cancer vs. HIV
2. **Cancer in People with HIV**
3. Opportunities to Enhance the Reach of HIV Service Delivery by Addressing Cancer in PWH
4. Conclusions/Invitation

# The complex and intersecting social, structural, behavioral, and lifestyle factors that affect cancer risk for PWH



# Cancer in People Living with HIV (PLWH)

Age-standardized mortality rates in PLWH and HIV-uninfected patients with cancer



PLWH have **higher cancer-specific mortality** than HIV-uninfected patients

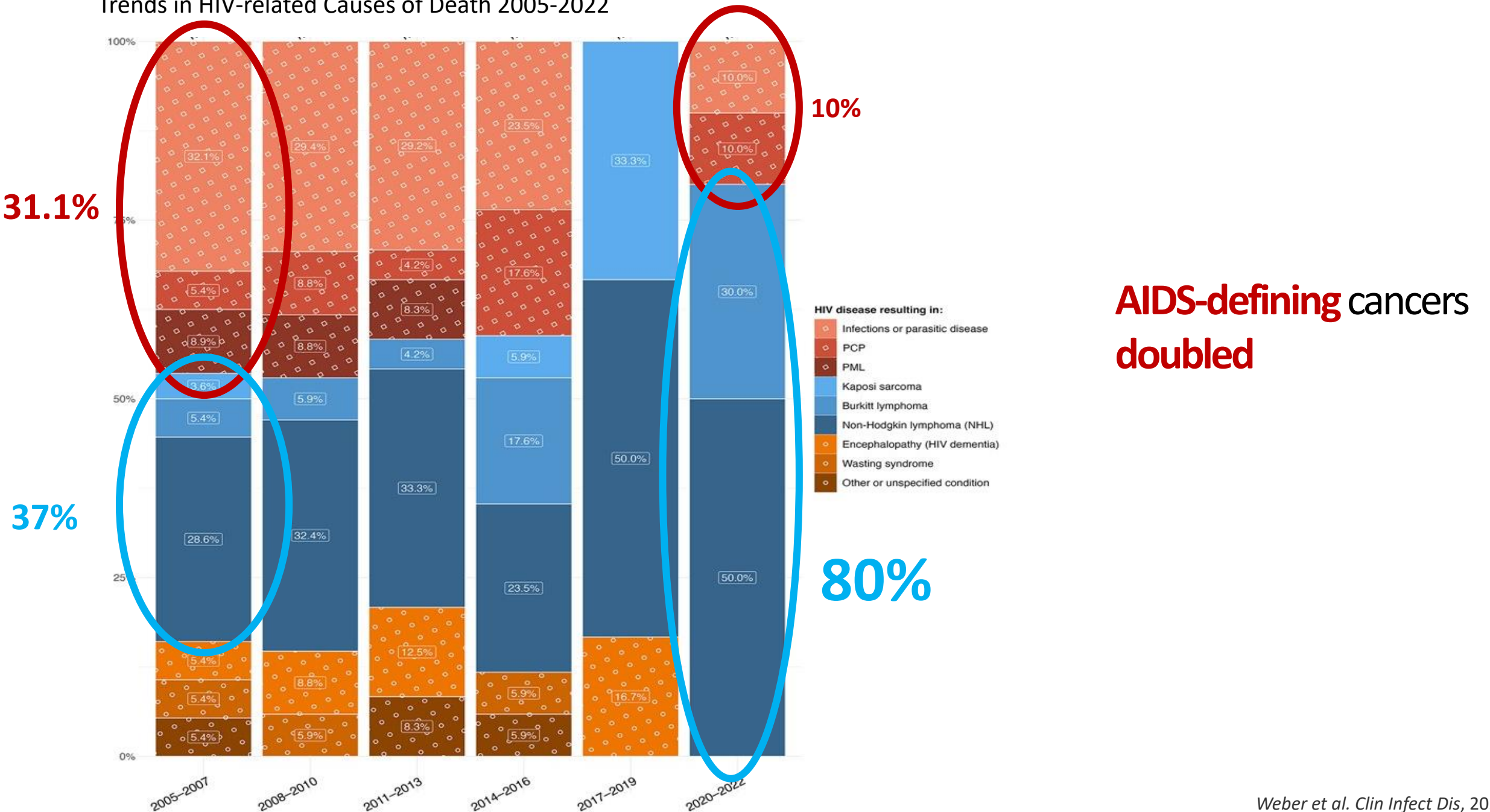
# Causes of death in PWH from the Swiss Cohort Study 2005-2022



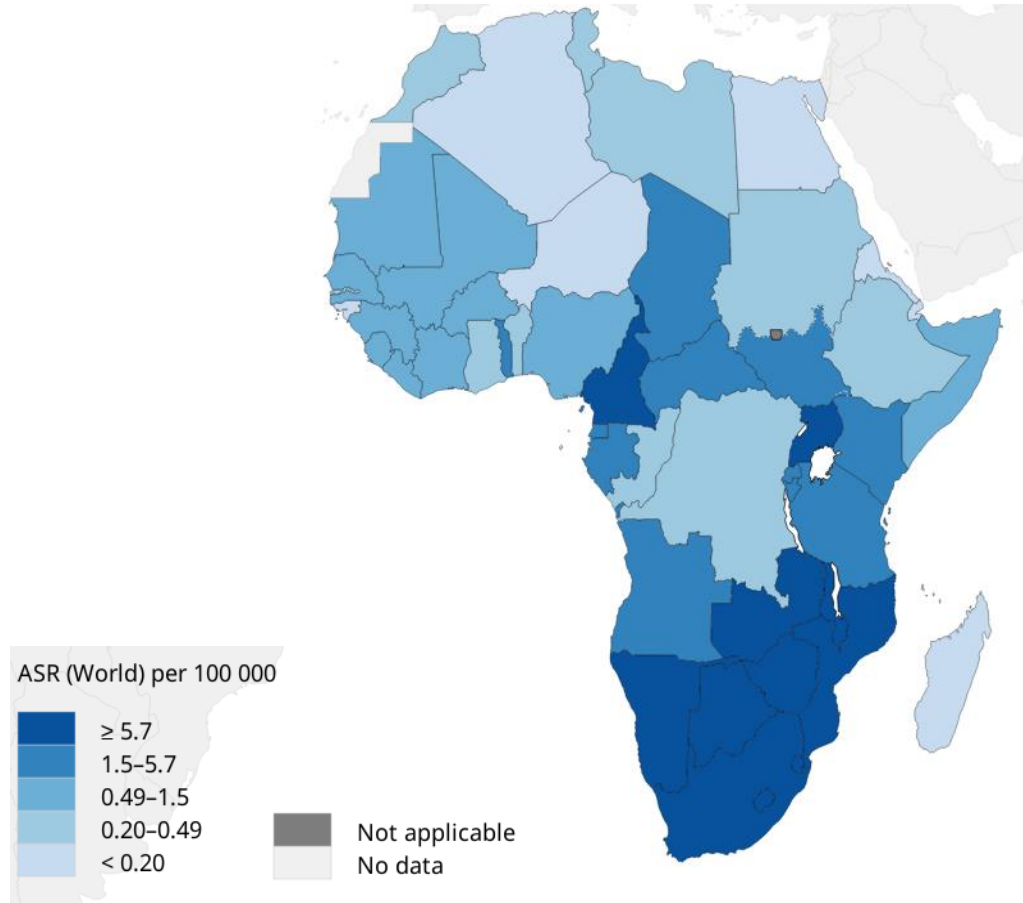
Non-AIDS defining cancers **doubled** as a **cause of death**



# Trends in HIV-related Causes of Death 2005-2022



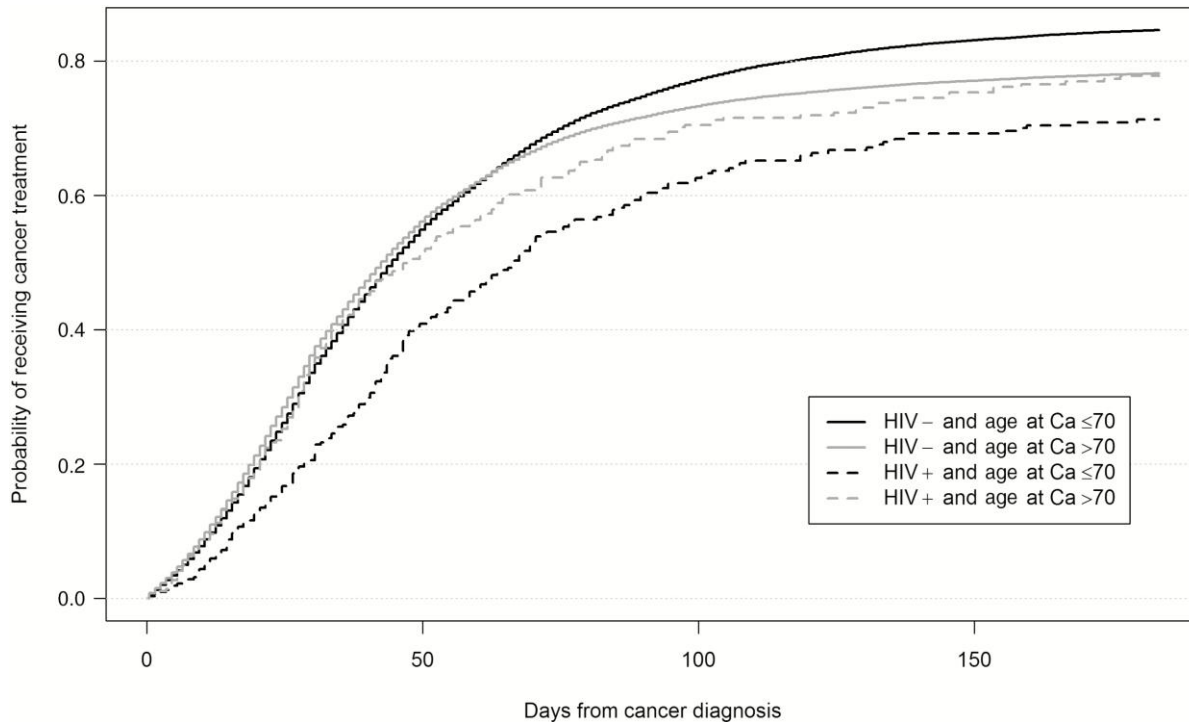
# Incidence rates of Kaposi sarcoma in 2020



GLOBOCAN 2022

# Cancer in People with HIV (PWH)

Probability of receiving cancer (Ca) treatment within 6 months of diagnosis, by age and HIV status.



PLWH experience **more barriers** to cancer care

- More likely to be diagnosed with cancer at advanced stage
- Less likely to receive cancer treatment
- Less likely to be enrolled in cancer clinical trials

# OUTLINE

1. Global Burden of Cancer vs. HIV
2. Cancer in People with HIV
3. **Opportunities to Enhance the Reach of HIV Service Delivery by Addressing Cancer in PWH**
4. Conclusion/Invitation

# Exemplars of HIV implementation interventions and strategies and potential adaptations for cancer in PWH

<b>Intervention in HIV</b>	<b>Potential Adaptation for Cancer in PWH</b>
<b>Single Tablet Regimens (STRs)</b>	Simplify cancer treatment dosing to improve adherence.
<b>Patient Navigation Services</b>	Provide navigation services to help cancer patients manage healthcare and treatment.
<b>Telemedicine and Digital Health</b>	Expand access to cancer care through telemedicine and digital platforms.
<b>Prevention and Screening Programs</b>	Develop widespread screening and prevention programs for high-risk populations.
<b>Integrated Care Models</b>	Implement multidisciplinary care models for complex cancer treatments.
<b>Community-Based Interventions</b>	Adapt outreach and education programs for cancer prevention and early detection.



# Simplifying cancer treatment dosing to improve adherence in PWH

## Treatment of advanced AIDS-associated Kaposi sarcoma in resource-limited settings: a three-arm, open-label, randomised, non-inferiority trial

Susan E Krown, Carlee B Moser, Patrick MacPhail, Roy M Matining, Catherine Godfrey, Stephanie R Caruso, Mina CHosseini-pour, Wadzanai Samaneka, Mulinda Nyirenda, Naftali W Busakhala, Fred M Okuku, Josphat Kosgei, Brenda Hoagland, Noluthando Mwelase, Vincent O Oliver, Henriette Burger, Rosie Mngqibisa, Mostafa Nokta, Thomas B Campbell, Margaret Z Borok, for the A5263/AMC066 protocol team

### Summary

**Background** Optimal treatment regimens for AIDS-associated Kaposi sarcoma, a frequent contributor to morbidity and mortality among people with HIV, have not been systematically evaluated in low-income and middle-income countries, where the disease is most common. In this study, we aimed to investigate optimal treatment strategies for advanced stage disease in areas of high prevalence and limited resources.

**Methods** In this open-label, non-inferiority trial, we enrolled people with HIV and advanced stage AIDS-associated Kaposi sarcoma attending 11 AIDS Clinical Trials Group sites in Brazil, Kenya, Malawi, South Africa, Uganda, and Zimbabwe. Eligible participants were randomly assigned (1:1:1) with a centralised computer system to receive either intravenous bleomycin and vincristine or oral etoposide (the investigational arms), or intravenous paclitaxel (the control arm), together with antiretroviral therapy (ART; combined efavirenz, tenofovir disoproxil fumarate, and emtricitabine). The primary outcome was progression-free survival (PFS) at week 48, using a 15% non-inferiority margin to compare the investigational groups against the active control group. Safety was assessed in all eligible treated study participants. The study was registered with ClinicalTrials.gov, NCT01435018.

**Findings** 334 participants were enrolled between Oct 1, 2013, and March 8, 2018, when the study was closed early due to inferiority of the bleomycin and vincristine plus ART arm, as per the recommendations of the Data and Safety Monitoring Board (DSMB). The etoposide plus ART arm also closed due to inferiority in March, 2016, following a DSMB recommendation. Week-48 PFS rates were higher in the paclitaxel plus ART arm than in both investigational arms. The absolute differences in PFS were -30% (95% CI -52 to -8) for the comparison of paclitaxel plus ART (week 48 PFS 50%, 32 to 67; n=59) and etoposide plus ART (20%, 6 to 33; n=59), and -20% (-33% to -7%) for the comparison of paclitaxel plus ART (64%, 55 to 73; n=138) and bleomycin and vincristine plus ART (44%, 35 to 53; n=132). Both CIs overlapped the non-inferiority margin. The most common adverse events, in 329 eligible participants who began treatment, were neutropenia (48 [15%]), low serum albumin (33 [10%]), weight loss (29 [9%]), and anaemia (28 [9%]), occurring at similar frequency across treatment arms.

**Interpretation** Non-inferiority of either investigational intervention was not shown, with paclitaxel plus ART showing superiority to both oral etoposide plus ART and bleomycin and vincristine plus ART, supporting its use in treating advanced AIDS-associated Kaposi sarcoma in resource-limited settings.

**Single agent** paclitaxel **superior** to combined bleomycin + vincristine for **HIV-associated Kaposi sarcoma** in LMIC

Could adapt HIV strategies that implemented and scaled single drug regimens

# Simplifying cancer treatment dosing to improve adherence in PWH



Blood 142 (2023) 4467–4469



The 65th ASH Annual Meeting Abstracts

## POSTER ABSTRACTS

### 626.AGGRESSIVE LYMPHOMAS: PROSPECTIVE THERAPEUTIC TRIALS

#### A Phase 1 Study of Subcutaneous Rituximab Hyaluronidase Combined with Local Standard-of-Care Chemotherapy for the Treatment of Diffuse Large B-Cell Lymphoma in Uganda

Henry Ddugnu, MD PhD<sup>1</sup>, Kelvin Mubiru, MS<sup>2</sup>, Scott V Adams, PhD<sup>3</sup>, Jacqueline Asea, MSN<sup>2</sup>, Joyce Kambugu, MBChB, MMED<sup>1</sup>, Rosemary Namagembe, BSN<sup>2</sup>, Prossy Namuli, BSN<sup>2</sup>, Jackson Orem, MBChB, PhD<sup>1</sup>, Camille Puronen, MDMPH<sup>4</sup>, Thomas S. Uldrick, MDMS<sup>3</sup>, Edus H Warren, MD PhD<sup>5,6</sup>, Manoj P Menon, MD<sup>5,6</sup>

<sup>1</sup>Uganda Cancer Institute, Kampala, Uganda

<sup>2</sup>Hutchinson Center Research Institute of Uganda., Kampala, Uganda

<sup>3</sup>Fred Hutchinson Cancer Center, Seattle

<sup>4</sup>University of Washington, Seattle

<sup>5</sup>Fred Hutchinson Cancer Center, Seattle, WA

<sup>6</sup>Division of Hematology and Oncology, University of Washington, Seattle, WA

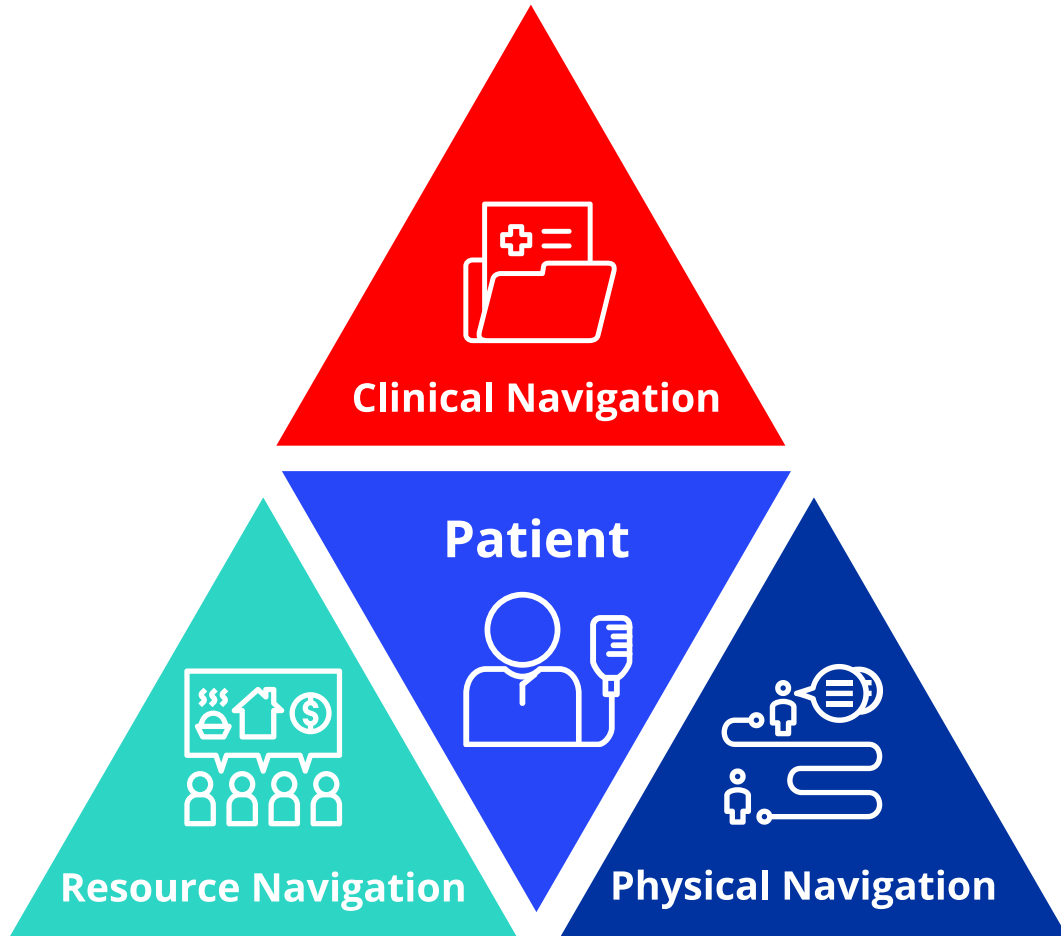
**Background:** Patients with Diffuse Large B-Cell Lymphoma (DLBCL) who are treated in low-resource settings have inferior outcomes compared to those in high-resource settings, with 12-month overall survival (OS) rates typically under 50% with standard chemotherapy (Gopal et al. [PMID:26934054]). Rituximab, an anti-CD20 monoclonal antibody, when combined with chemotherapy, improves OS for DLBCL and is the standard of care for patients with CD20+ DLBCL in resource-rich settings. Despite receiving FDA approval over 25 years ago and its subsequent inclusion in the World Health Organization's List of Essential Medicines, Rituximab is not readily available or routinely used in sub-Saharan Africa (SSA). This is in part due to

**Subcutaneous** rituximab is safe and efficacious and **precludes need for IV infusions** for **Lymphoma** treatment in **Uganda**

Could adapt HIV strategies that are implementing injectable PrEP and ART

# Patient Navigation Services

American Cancer Society Patient Navigation Program in Kenya



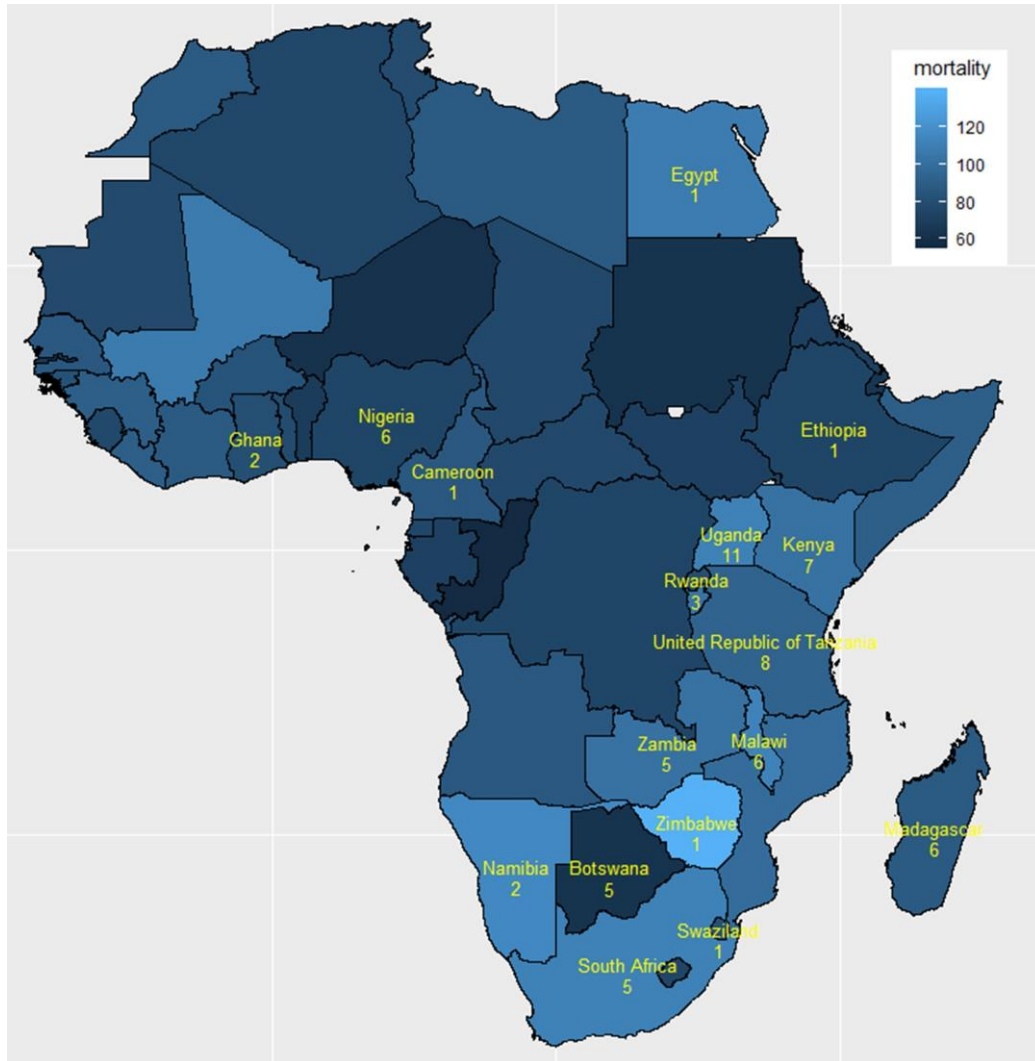
The American Cancer Society BEACON Initiative supports health systems in LMICs to design, implement, and sustain **cancer patient navigation** programs.

Could adapt HIV strategies for engagement in care that have implemented and scaled patient navigation



# Telemedicine and Digital Health

## Digital Health Pilots in Cancer Programs



Most digital health programs for cancer are isolated pilots

Reach should be extended to PWH with cancer by adapting from successful HIV digital health programs and leveraging impl sci to address barriers to spread.

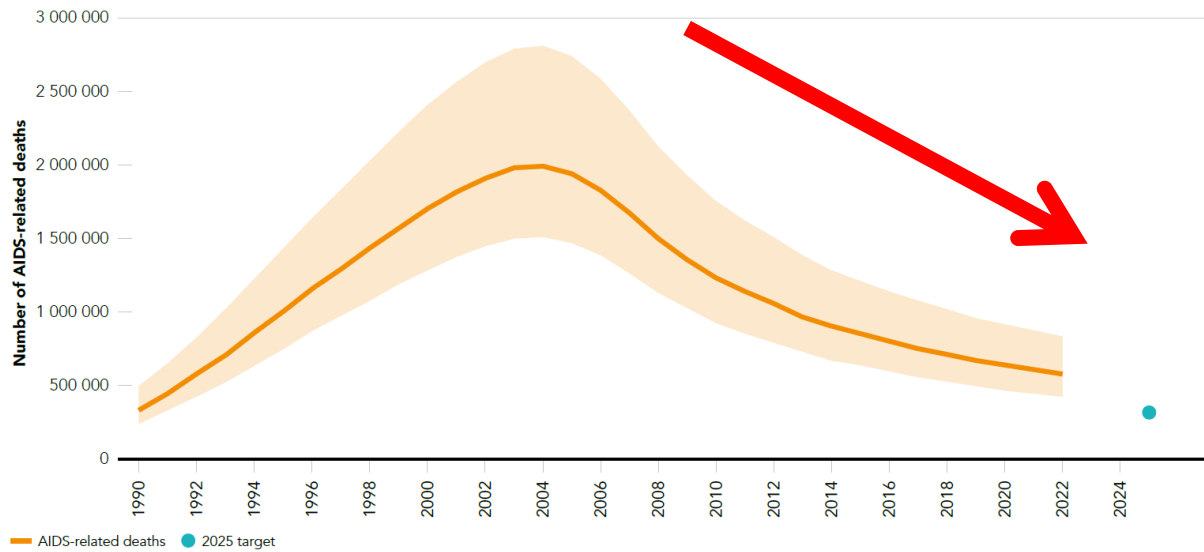
# OUTLINE

1. Global Burden of Cancer vs. HIV
2. Cancer in People with HIV
3. Opportunities to Enhance the Reach of HIV Service Delivery by Addressing Cancer in PWH
- 4. Conclusion/Invitation**

# 1. Cancer-related deaths are on the rise

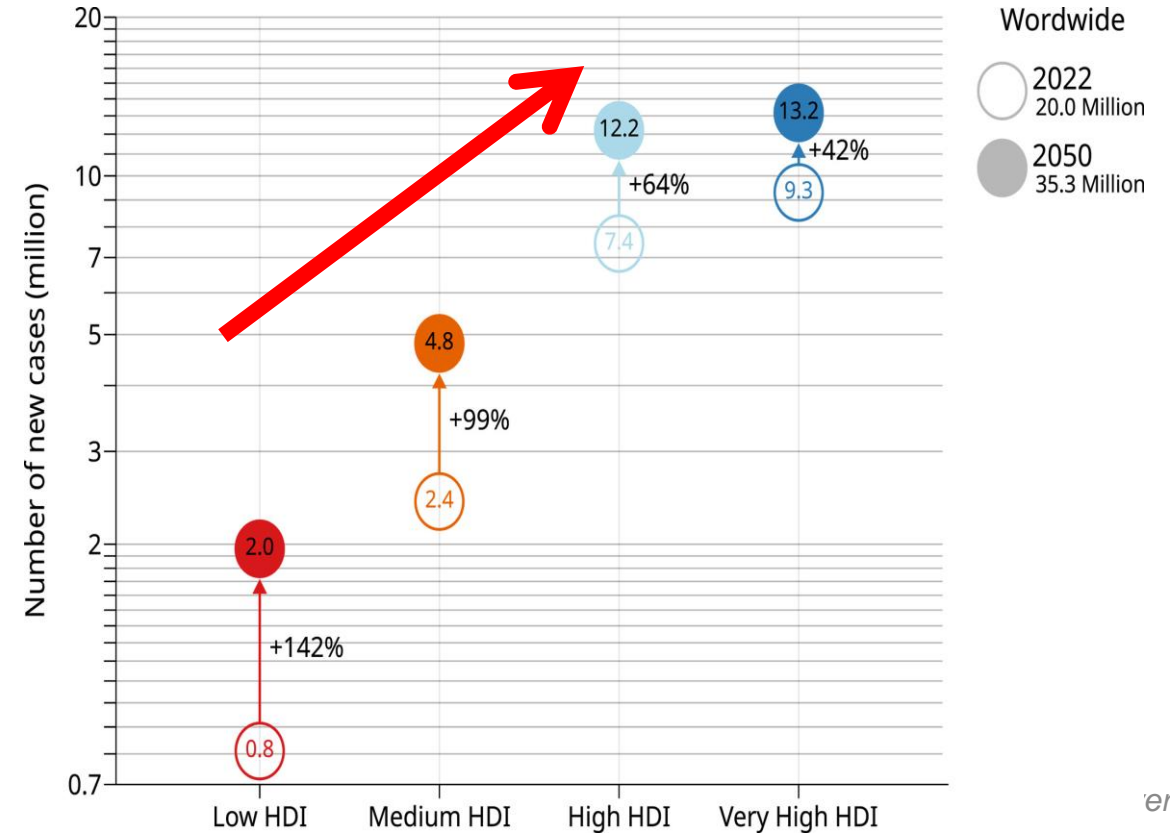
## HIV-related deaths

Figure 12.2 Number of AIDS-related deaths, global, 1990–2022, and 2025 target



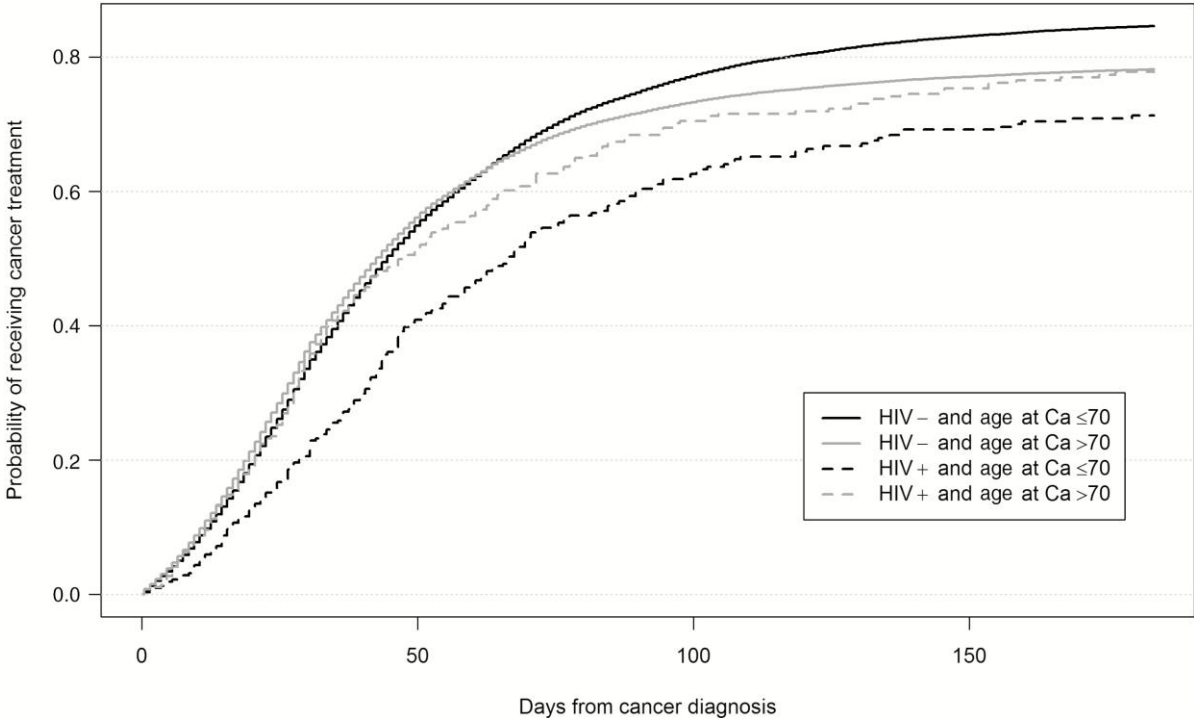
Source: UNAIDS epidemiological estimates, 2023 (<https://aidsinfo.unaids.org/>).

## Cancer-related deaths



# 2. PWH Get More Cancer and Experience More Barriers

Probability of receiving cancer (Ca) treatment within 6 months of diagnosis, by age and HIV status.

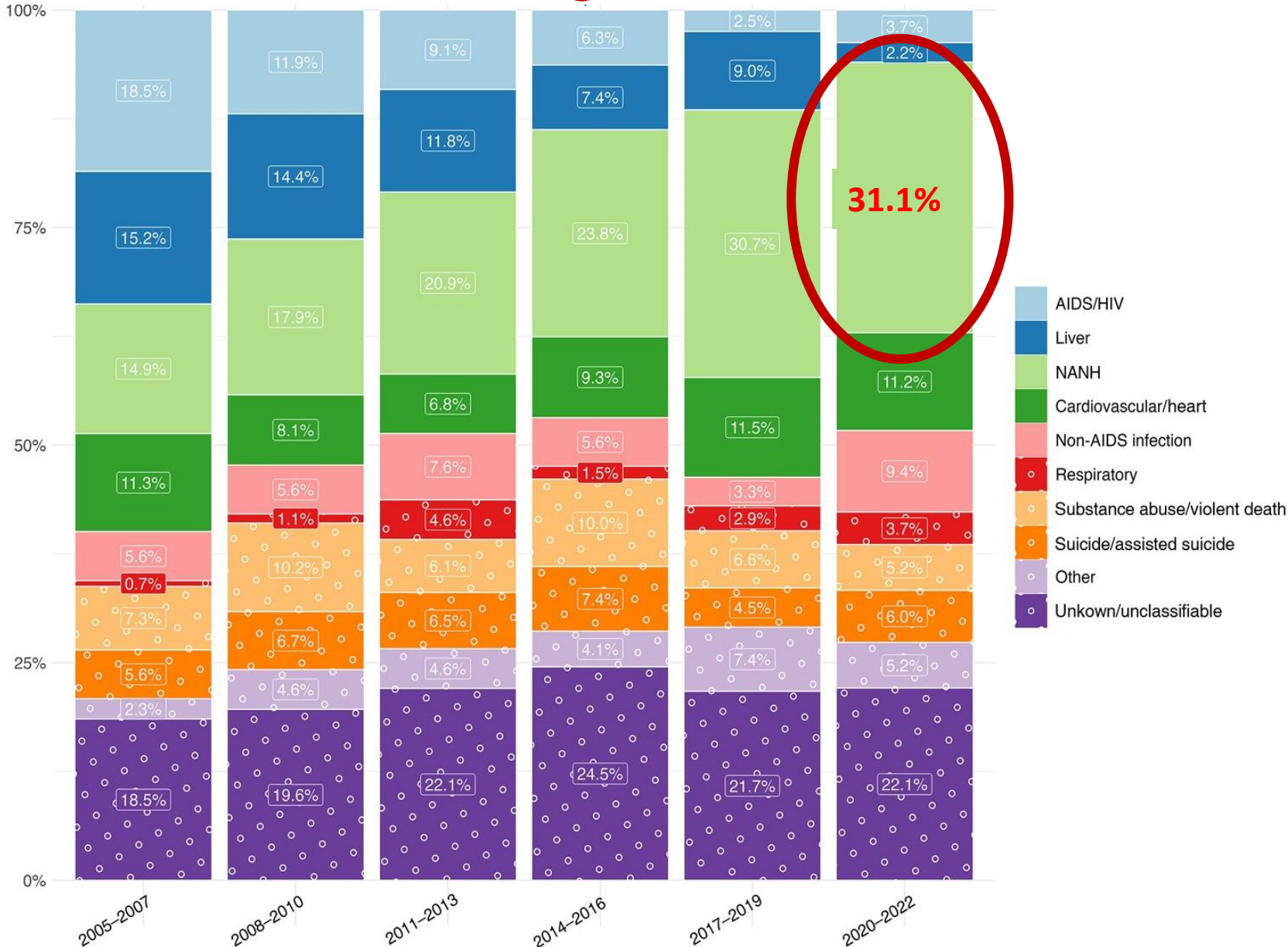


PWH are more likely to get cancer

PWH with cancer experience **more barriers** to cancer care

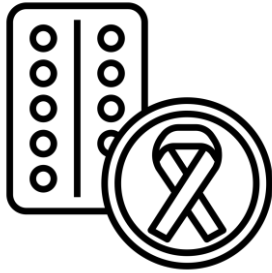
Rositch AF et al. Clin Infect Dis. 2018

### 3. Cancer is a Leading Cause of Death in PWH

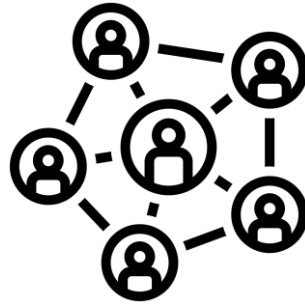


Cancer is a leading cause of death in PWH

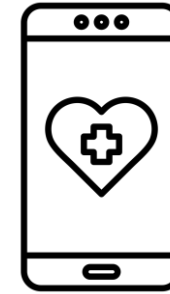
## 4. Opportunities Exist for Successful HIV implementation strategies to be adapted to extend the reach of services for PWH with cancer



**Simplifying cancer treatment for  
PWH**



**Patient navigation services for PWH  
with cancer**



**Digital Health and Telemedicine**

# Thank you!

## Washington University

- Elvin Geng
- Betsy Abente

## Kenya Medical Research Institute (KEMRI)

- Elizabeth Bukusi
- Eliud Akama
- Fridah Adhiambo



Washington University in St. Louis

SCHOOL OF MEDICINE