

ACHIEVING 90:90:90:

A GLOBAL GAME CHANGER FOR PUBLIC HEALTH

Stefano Vella MD Istituto Superiore di Sanità - Rome - Italy



ACHIEVING 90:90:90: A GLOBAL GAME CHANGER FOR PUBLIC HEALTH

The Goal

The Tools

Addressing the barriers

From Durban to Durban

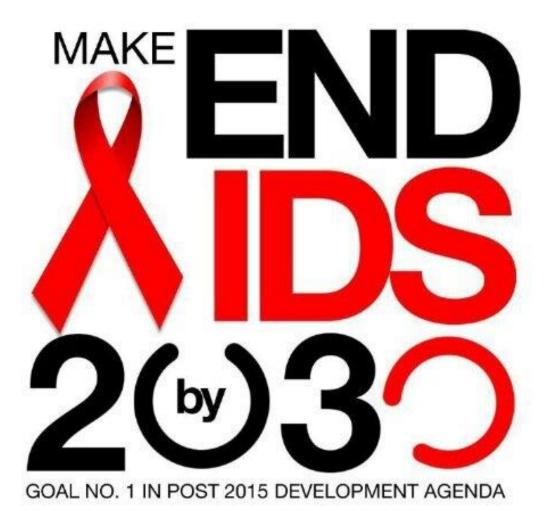
Open Working Group proposal for

Sustainable Development Goals



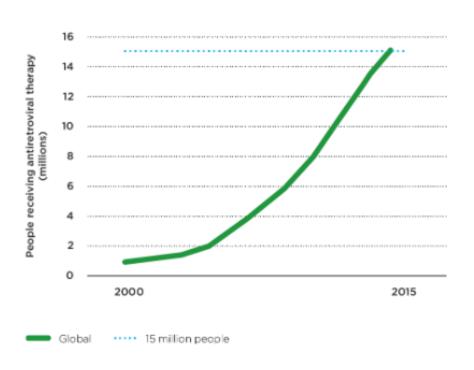
Goal #3 - Ensure healthy lives and promote well-being for all at all ages

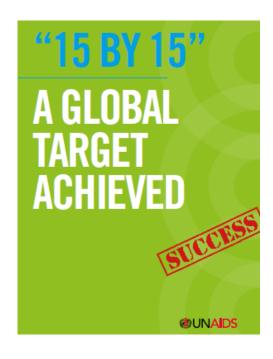
- ✓ By 2030, reduce the global maternal mortality ratio to less than 70 per 100,000 live births
 - ✓ By 2030, end preventable deaths of newborns and children under 5 years of age
 - ✓ By 2030, end the epidemics of AIDS, tuberculosis, malaria and neglected tropical diseases and combat hepatitis, water-borne diseases and other communicable diseases
 - ✓ By 2030, reduce by one third premature mortality from non-communicable diseases
 - ✓ By 2030, through prevention and treatment promote mental health and well being
 - **✓** By 2030, ensure universal access to sexual and reproductive health-care services.



2015 an amazing target achieved

Number of people receiving antiretroviral therapy, 2000–2015







GLOBAL MORTALITY AND INCIDENCE DECREASED....

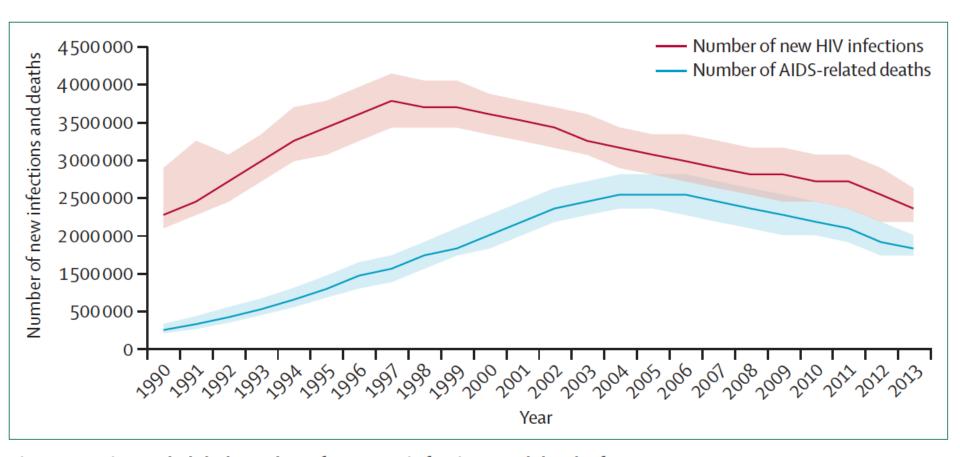
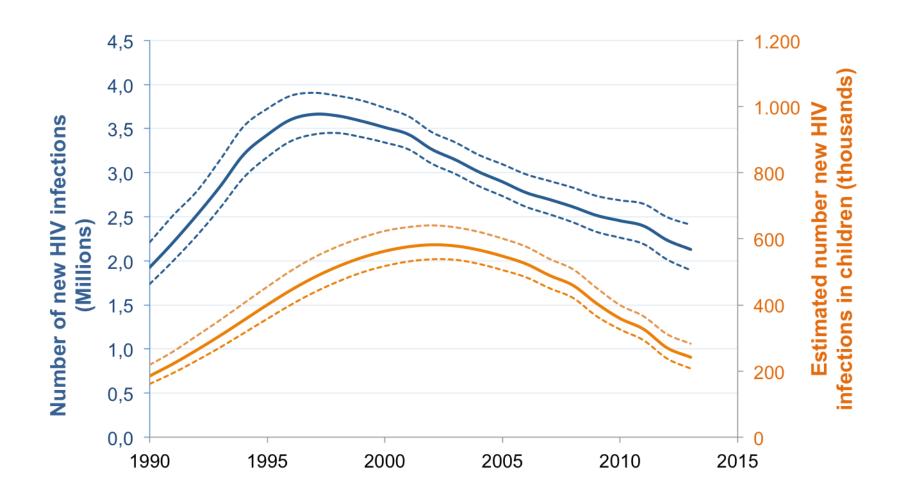


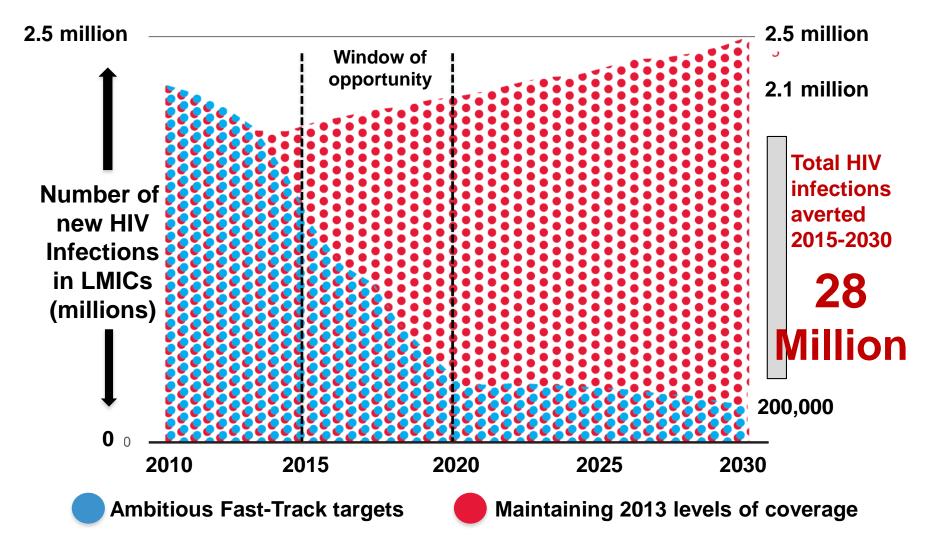
Figure 1: Estimated global number of new HIV infections and deaths from AIDS, 1990–2013 Source: UNAIDS 2013 global fact sheet.¹³ Shaded areas indicate uncertainty bounds.

...however, the global number of new HIV infections in adults and children is still unacceptably high



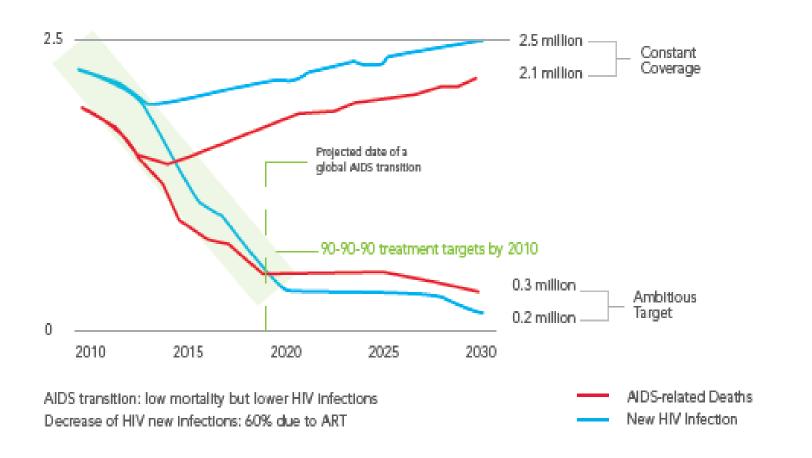
Source: UNAIDS Global Report 2014

"The AIDS response is at a crucial juncture, both in its immediate trajectory and its sustainability..."



Source: Adapted from UNAIDS Fast-track Report

For clear, HIV epidemic will rebound without change in coverage by 2020



ACHIEVING 90:90:90: A GLOBAL GAME CHANGER FOR PUBLIC HEALTH

The Goal

The Tools

Addressing the barriers

From Durban to Durban



Needle **Exchange**

Drucker E, AIDS 1998



Auvert B, PloS Med 2005 Gray R, Lancet 2007 Bailey R, Lancet 2007





Grosskurth H, Lancet 2000







Abdool Karim Q, Science 2010

HIV **PREVENTION**

Male Condoms



Oral pre-exposure prophylaxis

Grant R, NEJM 2010 (MSM) Baeten J, NEJM 2012 (Couples) Thigpen M, NEJM 2012 (Heterosexuals) Choopanya K, Lancet 2013 (IDU)

HIV Counselling and Testing

Coates T, Lancet 2000



Sweat M, Lancet 2011



Opioid substitution therapy

Mathers BM, Lancet 2010

Treatment for prevention

Cohen M, NEJM, 2011 Donnell D, Lancet 2010 Tanser, Science 2013

Behavioural Intervention



Note: PMTCT, Screening transfusions, Universal precautions, etc. have not been included

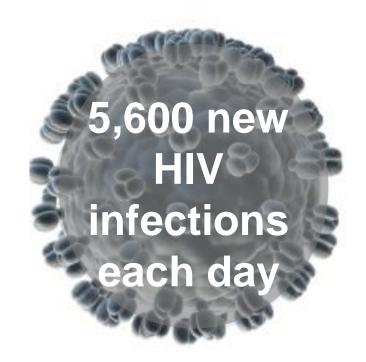
Despite impressive progress with prevention, the spread of HIV has yet to be controlled!

In 2014, worldwide there were:

1.2 million HIV deaths

36.9 million living with HIV

2.0 million new infections





Source: UNAIDS Global Report 2014

Stigma: Major impediment to HIV prevention and treatment



Stigma impedes AIDS prevention

nature

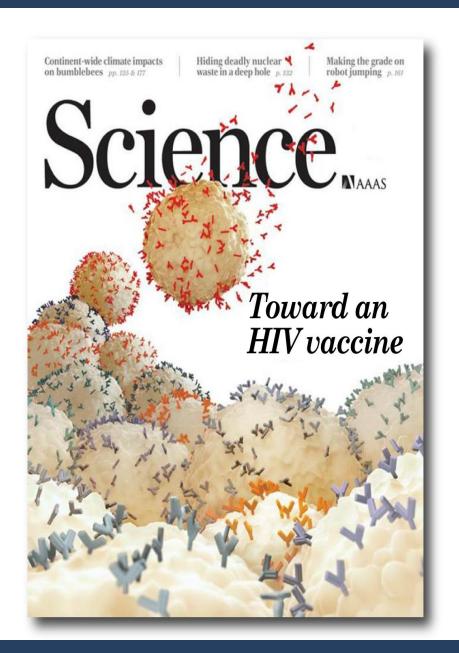
Medical advances cannot help those who deny they are at risk of HIV and avoid HIV tests. **Salim S. Abdool Karim** describes how such attitudes may be overcome.



Source: UNAIDS Together we will end AIDS 2012

New Prevention Technologies gives hope

- PrEP works (when used)
 - New meds and dosing regimens for oral PrEP may improve uptake, \u00c4cost
- Circumcision is highly effective
- Microbicides
 - Rectal gels may offer new anal protection
 - Rings may offer MPT opportunities
- Harm reduction for IDU works, and shall be implemented
- Vaccine may be closer than thought just a couple of years ago



Passive Transfer of Neutralizing Antibodies for Prevention of SHIV Infection



Protection of Macaques
against Vaginal
Transmission of a
Pathogenic HIV-1/SIV
Chimeric Virus by
Passive Infusion of
Neutralizing Antibodies

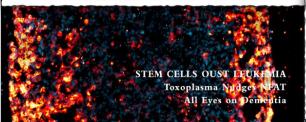
JR Mascola, SS Frankel, DL Birx, MG Lewis, et al.

Neutralization of Mir protects against septic snock
Antibody protection against mucosal HIV-1 transmission
A new amyloid β-protein catabolic pathway
HER-Z/neu inhibitor for cancer therapy



Passive Transfer of Modest
Titers of Potent and
Broadly Neutralizing
Anti-HIV Monoclonal
Antibodies Block SHIV
Infection in Macaques

M Shingai, DR Burton, MC Nussenzweig, MA Martin, Y Nishimura, et al.





Highly Potent HIV-Specific
Antibody Neutralization in
vitro Translates into
Effective Protection
against Mucosal SHIV
Challenge in vivo

B Moldt, DI Watkins, P Poignard, DR Burton, et al.

Published online June 18, 2015

Science

HIV-1 Neutralizing Antibodies Induced by Native-Like Envelope Trimers

RW Sanders, H Dean, DR Burton, JP Moore, et al.

Stabilized form of native trimer stimulated autologous NAb production in rabbits and non-human primates



June 22, 2015

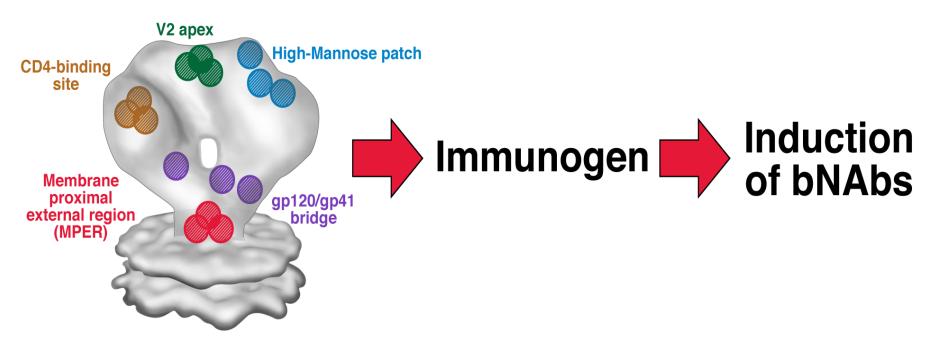
Crystal Structure, Conformational Fixation, and Entry-related Interactions of Mature Ligand-free HIV-1 Env

YD Kwon, M Pancera, P Acharya, IS Georgiev, JR Mascola, PD Kwong, et al.

- Unliganded HIV-1 envelope trimer, capable of binding broadly neutralizing antibodies
- Characterized and fixed in pre-fusion, closed conformation
- Potential utility as vaccine immunogen

 A. FAUCI, IAS 2015

Fundamental Challenge in HIV Vaccinology: Convert Neutralizing Epitopes to Immunogens Inducing bNAbs



Neutralizing epitopes

One day it may complement TASP However, an HIV vaccine is not there, yet.

•

So, lets focus on what we have

and which we know it works



Needle Exchange

Drucker E, AIDS 1998



Auvert B, PloS Med 2005 Gray R, Lancet 2007 Bailey R, Lancet 2007





Grosskurth H, Lancet 2000



Microbicides for women

Abdool Karim Q, Science 2010





PREVENTION

HIV





Truvada Truvad

Oral pre-exposure prophylaxis

Grant R, NEJM 2010 (MSM)
Baeten J, NEJM 2012 (Couples)
Thigpen M, NEJM 2012 (Heterosexuals)
Choopanya K, Lancet 2013 (IDU)

HIV Counselling and Testing

Coates T, Lancet 2000 Sweat M, Lancet 2011





Opioid substitution therapy

Mathers BM, Lancet 2010

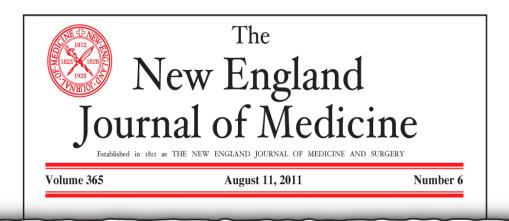
Treatment for prevention

Cohen M, NEJM, 2011 Donnell D, Lancet 2010 Tanser, Science 2013

Behavioural Intervention



Note: PMTCT, Screening transfusions, Universal precautions, etc. have not been included

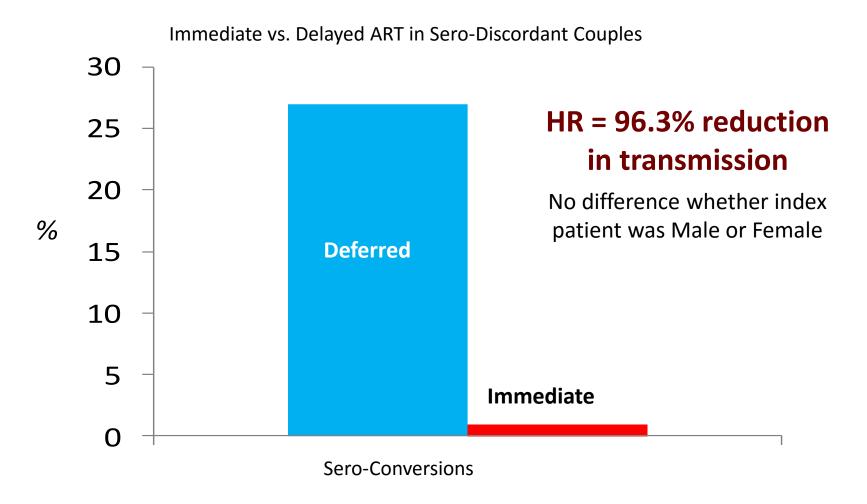


Prevention of HIV-1 Infection with Early Antiretroviral Therapy

HPTN 052 Study Team

- 1,763 HIV-serodiscordant couples in 9 countries
- 96% reduction in HIV transmission when ART started in HIV-infected partner at CD4 count of 350-550 compared to <250</p>

HPTN 052: treatment as prevention

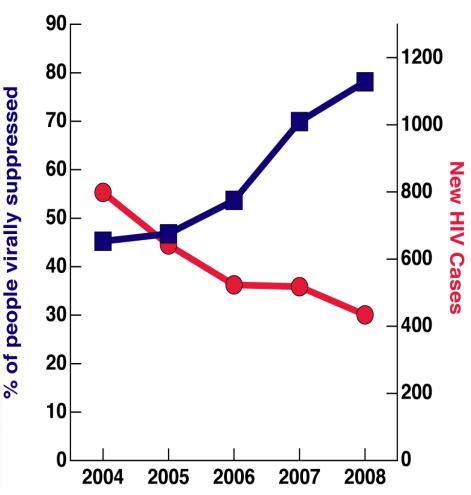


Treatment as Prevention in San Francisco



Decreases in Community Viral Load Are Accompanied by Reductions in New HIV Infections in San Francisco

M Das, PL Chu, GN Colfax, et al.



■ When viral suppression rose from 45 to 78%, HIV incidence fell by 45%

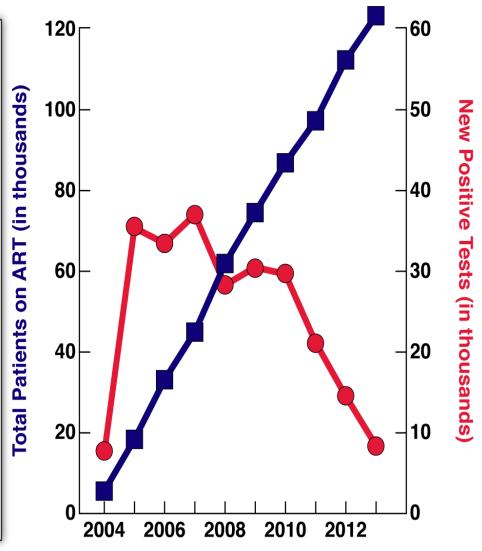
Treatment as Prevention in Rwanda



February 2015

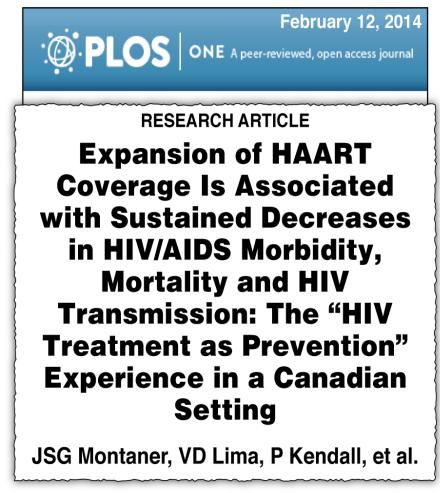
Nationwide Evaluation of Antiretroviral Therapy Coverage on Prevention in Rwanda: A Multisectional Time-Trend Analysis

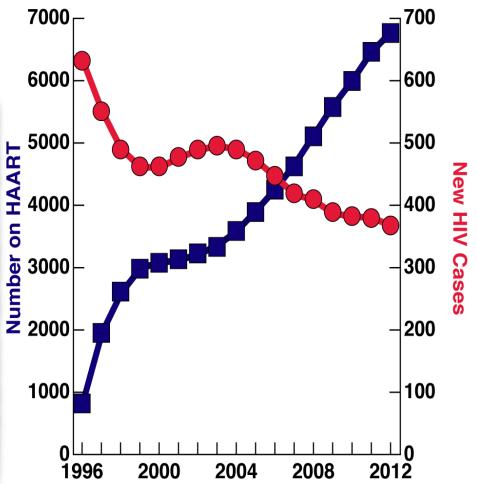
S Kanters, S Nsanzimana, T Barnighausen, JSG Montaner, et al.



Every 10% increase in ART coverage associated with a 6% decrease in HIV incidence

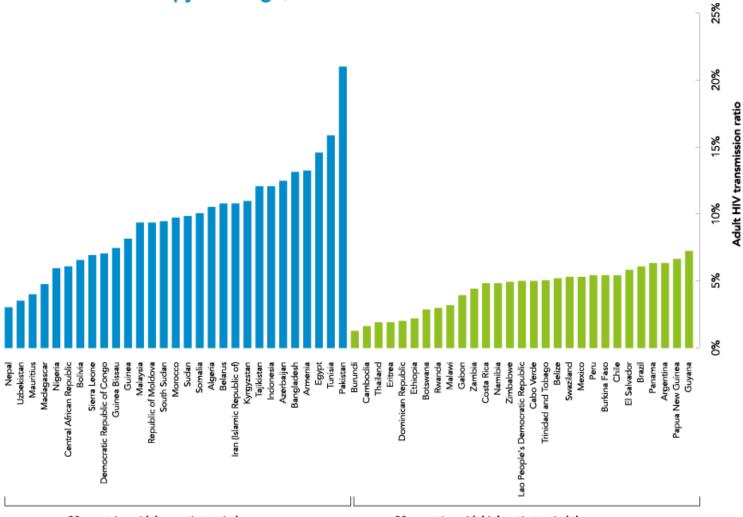
Treatment as Prevention in British Columbia





■ When ART coverage rose from 11 to 57%, HIV incidence fell by 42%

Adult HIV transmission rate in low- and middle-income countries with high and low antiretroviral therapy coverage, 2013



90 90 90: THE UNAIDS STRATEGY TO FURTHER CURB THE HIV EPIDEMIC

based on expanded access to treatment and on the "treatment as prevention" concept

90%

of all people living with HIV will know their HIV status 90%

of all people diagnosed with HIV will receive sustained antiretroviral therapy.

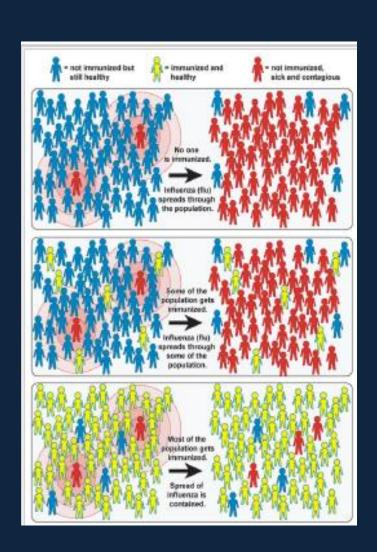
90%

of all people receiving antiretroviral therapy will have durable suppression.

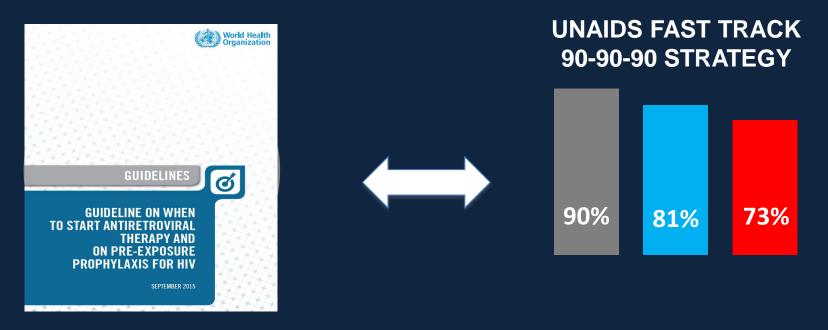
Zero new HIV infections. Zero discrimination. Zero AIDS-related deaths.



The "herd immunity" vaccine concept (which is usually applied to susceptibles to a virus) now applied to HIV transmitters



THE NEW WHO ELIGIBILITY CRITERIA AND THE UNAIDS 90 90 90 TARGETS ARE CONVERGING ELEMENTS OF THE SAME GOAL: ENDING AIDS BY 2030



By combining the personal health benefit (reducing HIV mortality & morbidity) with the Public Health benefit (reducing transmission) they foster universal access to care and treatment, provide operational advantage, and contribute to global equity

Wew eligibility criteria: treatme of CDA hew be started irrespective of Can be started irrespective.



GUIDELINE ON WHEN TO START ANTIRETROVIRAL THERAPY AND **ON PRE-EXPOSURE** PROPHYLAXIS FOR HIV

SEPTEMBER 2015

4.3.1 When to start ART in adults (>19 years old)

Recommendation

 ART should be initiated in all adults living with HIV regardless of WHO clinical stage and at any CD4 cell count (strong recommendation, moderate-quality evidence).

As a priority, ART should be initiated in all adults with severe or advanced HIV clinical disease (WHO clinical stage 3 or 4) and adults with CD4 count ≤350 cells/mm³ (strong recommendation, moderate-quality evidence).

Source:

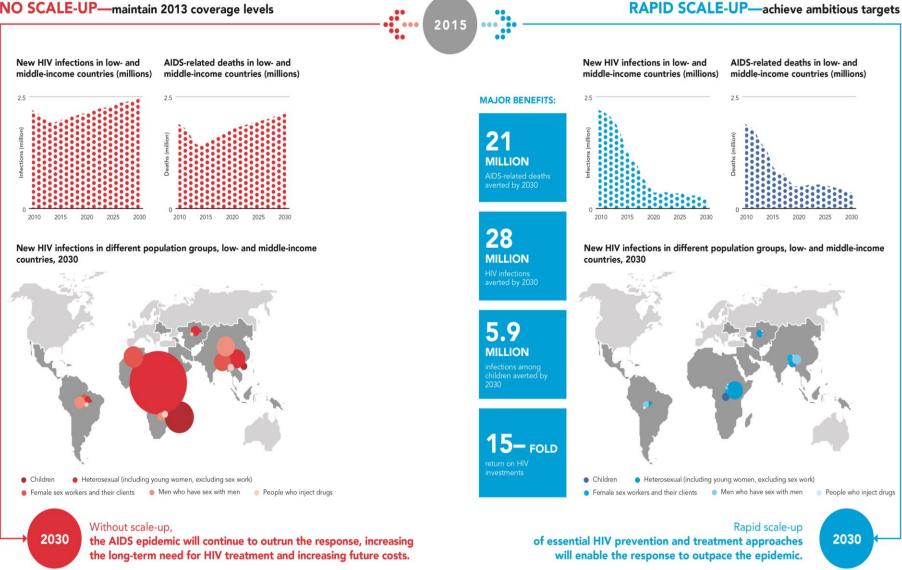
Guideline on when to start antiretroviral therapy and on pre-exposure prophylaxis for HIV Geneva: World Health Organization 2015.

(http://www.who.int/hiv/pub/guidelines/earlyrelease-arv/en/).

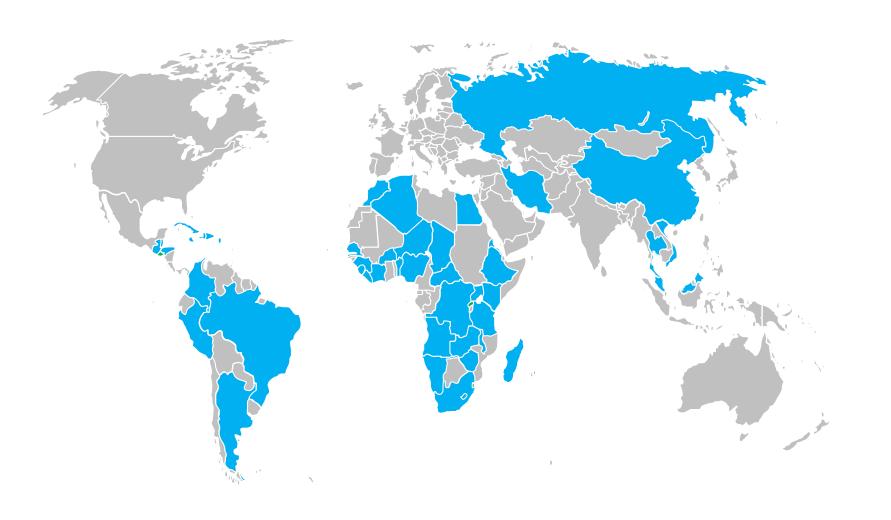
Consolidated guidelines on the use of antiretroviral drugs for treating and preventing HIV infection Recommendations for a public health approach Geneva: World Health Organization, 2013. (http://www.who.int/hiv/pub/guidelines/arv2013/download/en/)

The Fast-Track

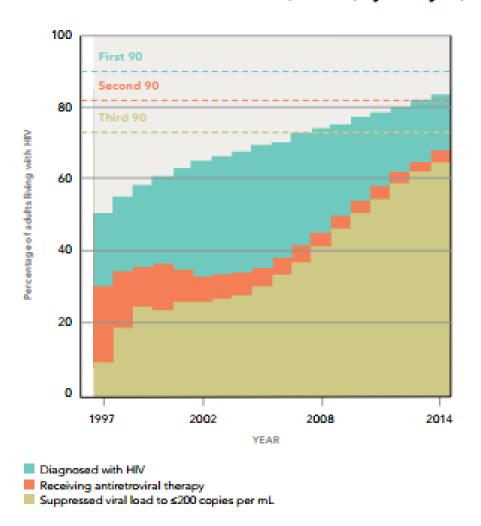
NO SCALE-UP—maintain 2013 coverage levels



Endorsement of the 90-90-90 treatment target

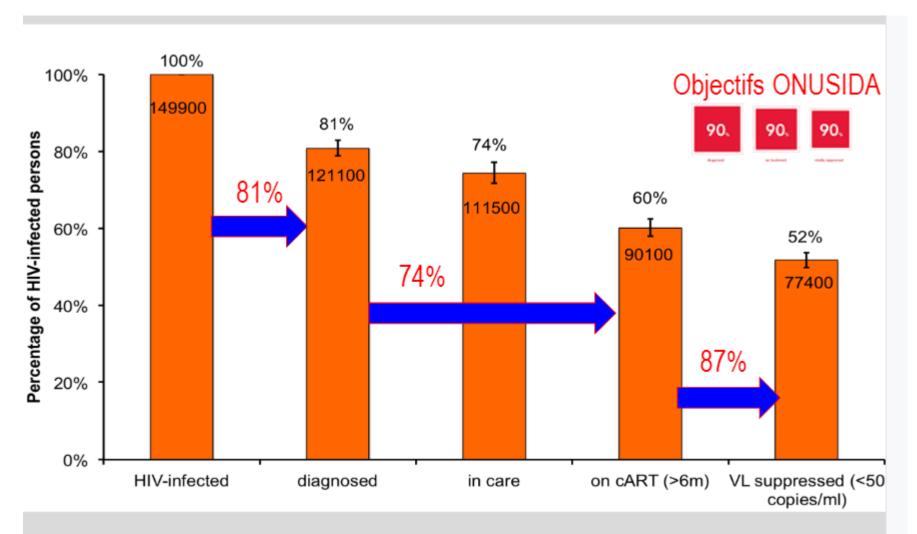


Cascade of care in British Columbia, Canada, by fiscal year, 1997-2014



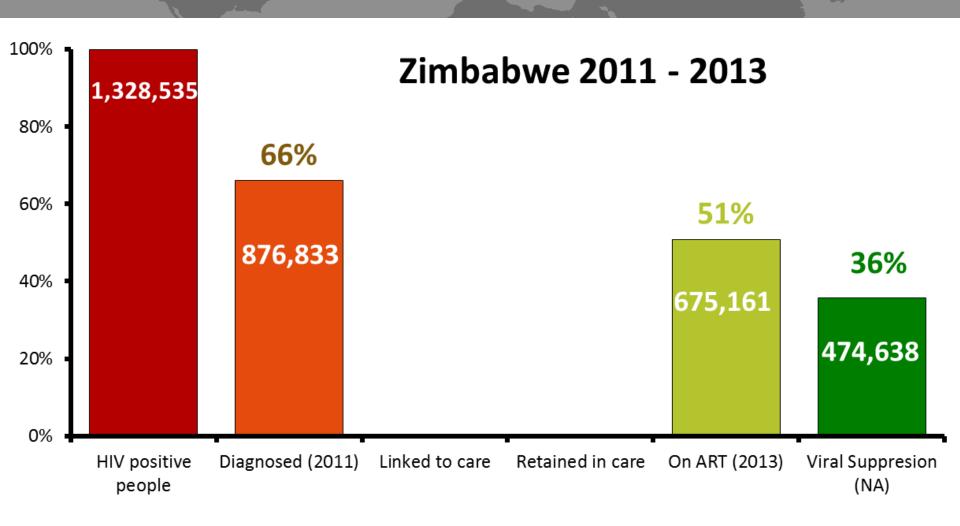
Source: British Columbia Centre for Excellence in HIV/AIDS Drug Treatment Program database (for antiretroviral use, viral load and CD4 count) and administrative data (ex. Medical Services Plan billings; hospitalization data from the Discharge Abstract Database).

FRANCE

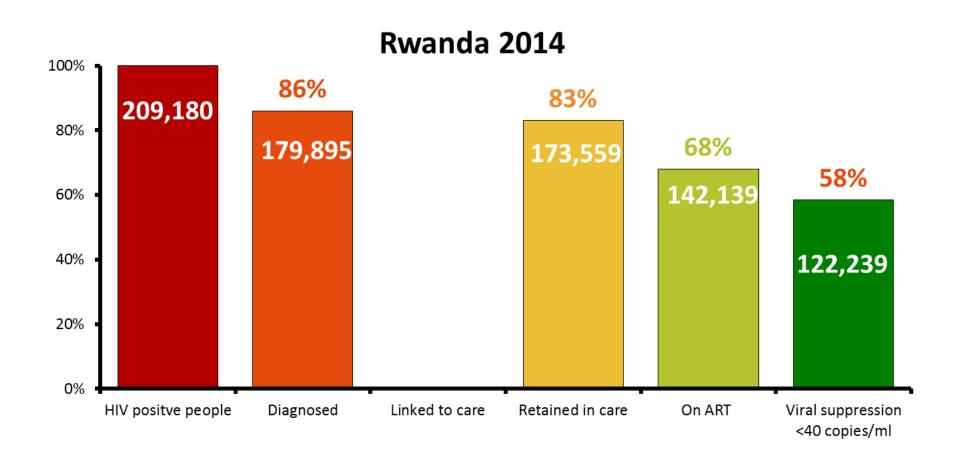


Supervie V. & Costagliola D. The spectrum of engagement in HIV care in France: strengths and gaps. 20th Conference on Retroviruses and Opportunistic Infections. Atlanta, USA: March 2013. Abstract #: 1030.

Cascade of HIV care – Zimbabwe 2014

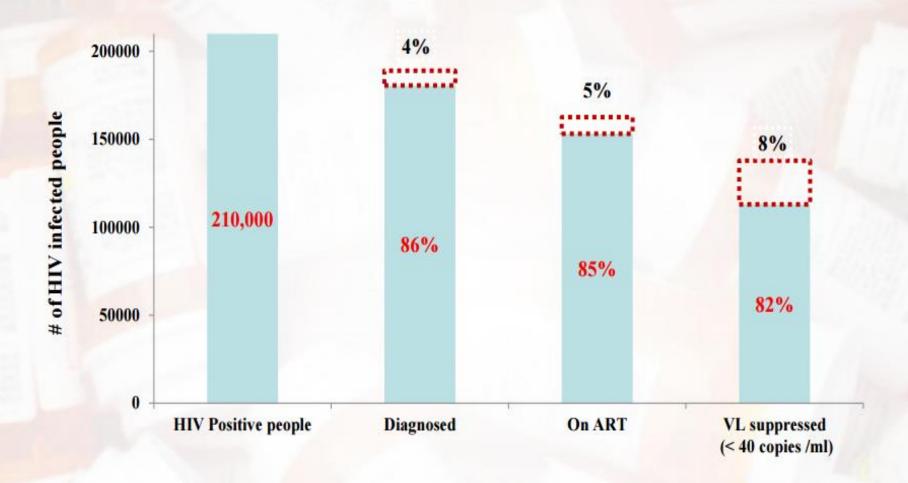


Cascade of HIV care – Rwanda 2014



Ref: Sabin Nsanzimana et al, IAS presentation 2015

Rwanda's Progress Towards 90-90-90



¹ Nsanzimana et al, The Lancet HIV feb 2015

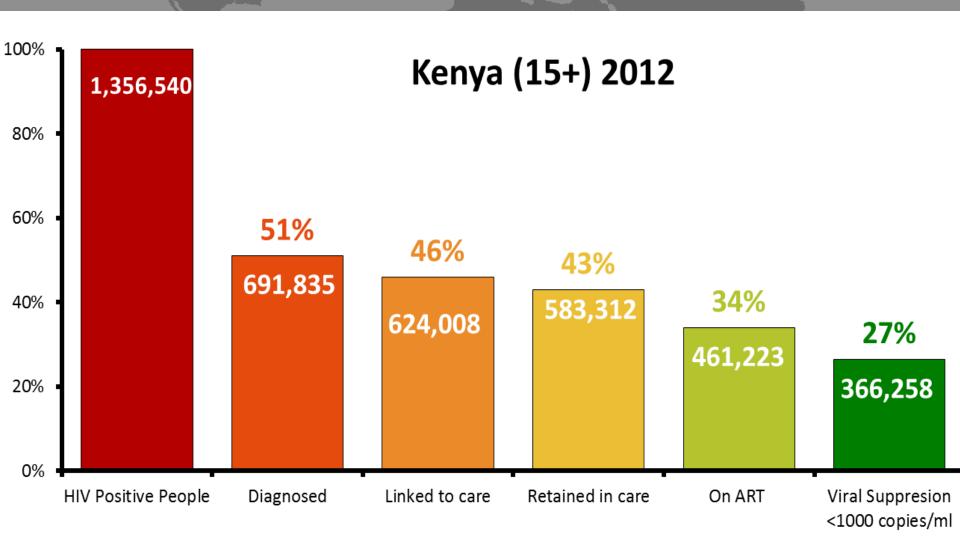




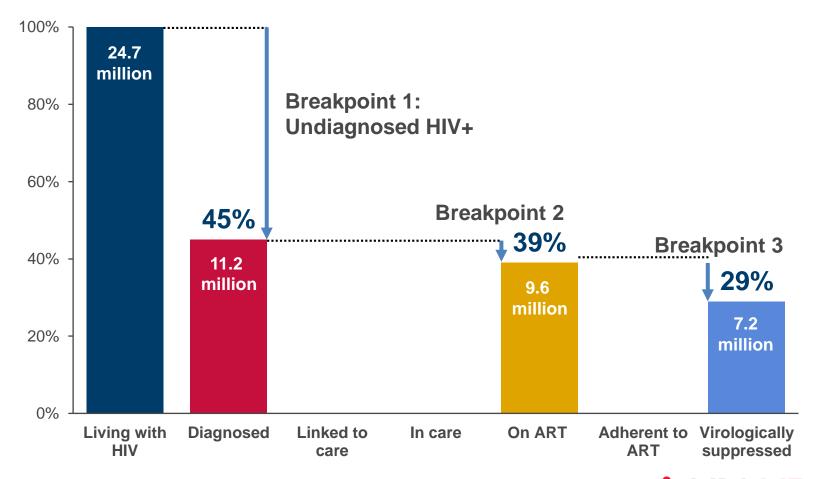
² Drug Resistance Monitoring in selected sites, 2013

³ HMIS, June 2015

Cascade of HIV care – Kenya 2012 (+15)



Cascade of HIV care - Sub-Saharan Africa



ACHIEVING 90:90:90: A GLOBAL GAME CHANGER FOR PUBLIC HEALTH

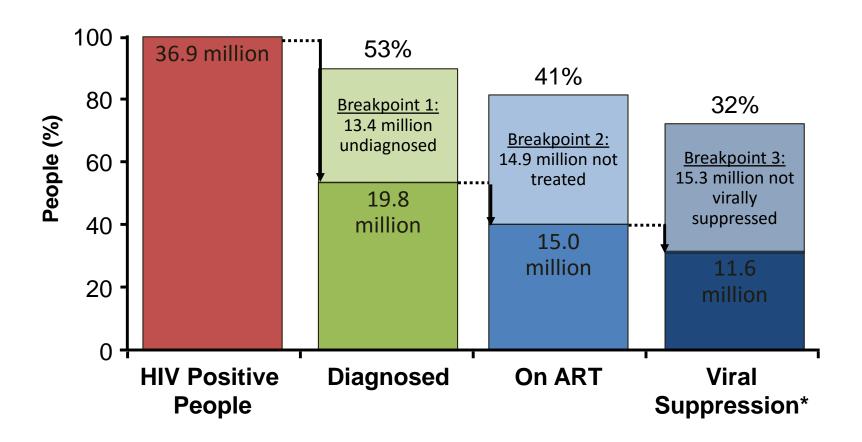
The Goal

The Tools

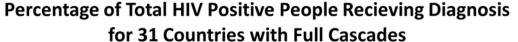
Addressing the barriers

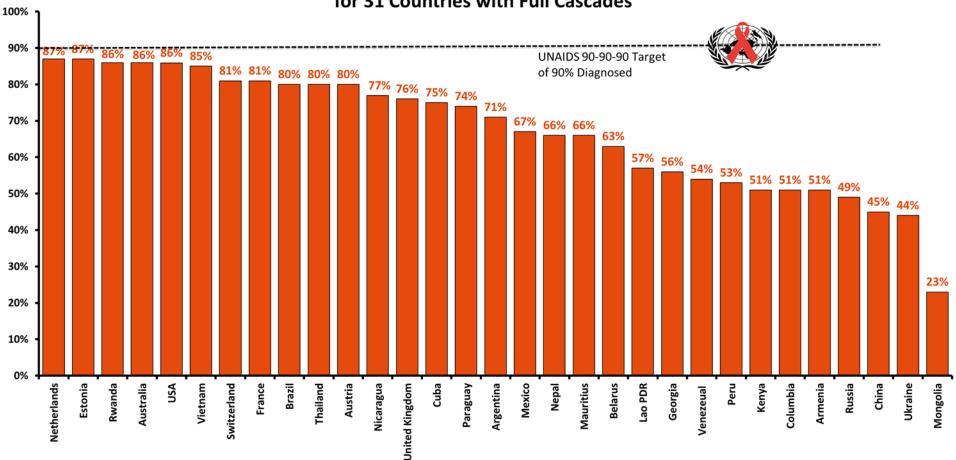
From Durban to Durban

UNAIDS 90-90-90: the gaps

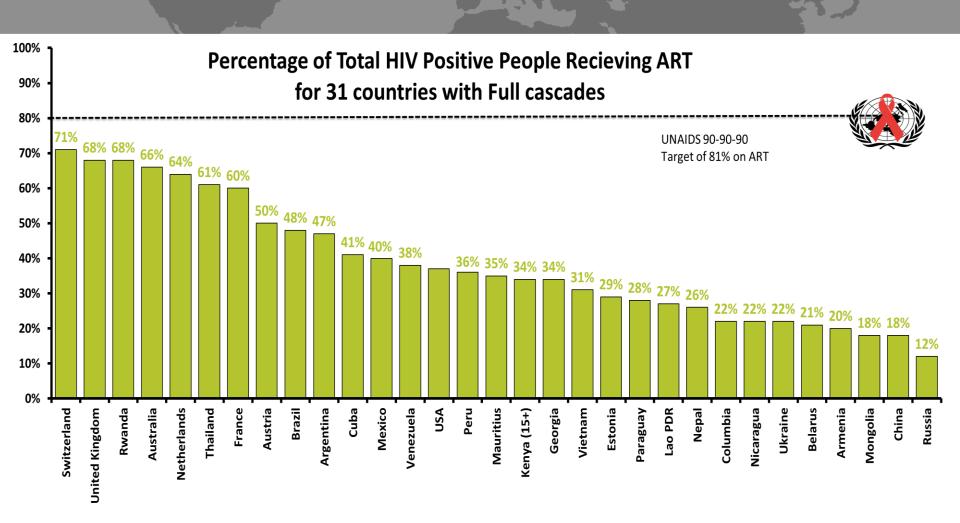


1st 90 – 90% diagnosis for Full cascades

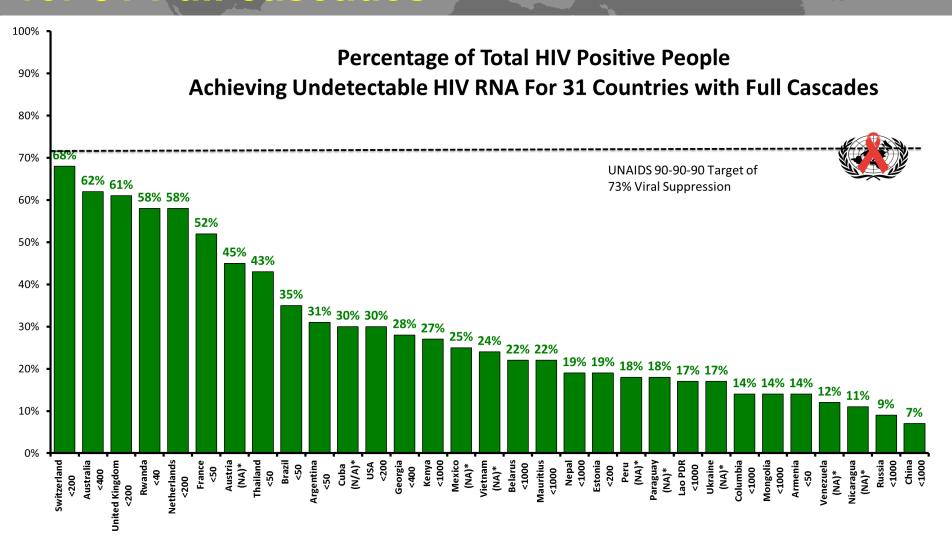




2nd 90 – 81% on ART for 31 Full cascades



3rd 90 – 73% achieving viral suppression for 31 Full cascades

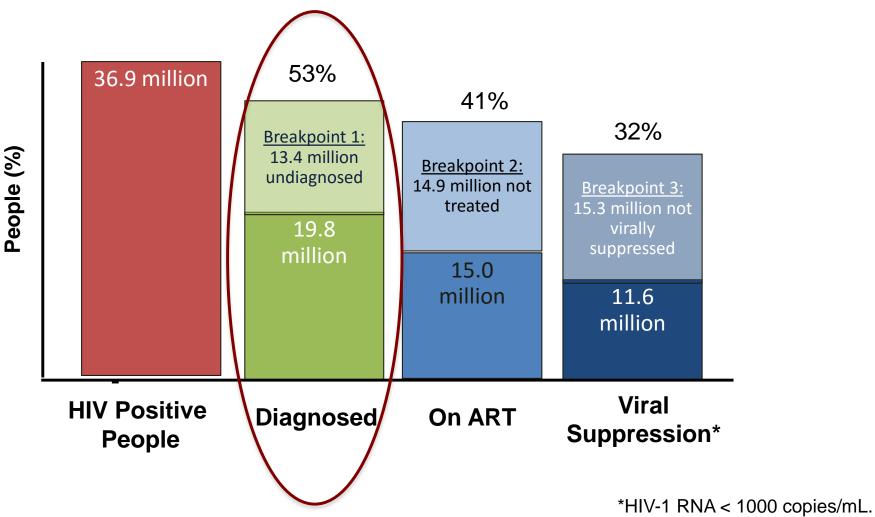


ACHIEVING 90:90:90: A GLOBAL GAME CHANGER FOR PUBLIC HEALTH

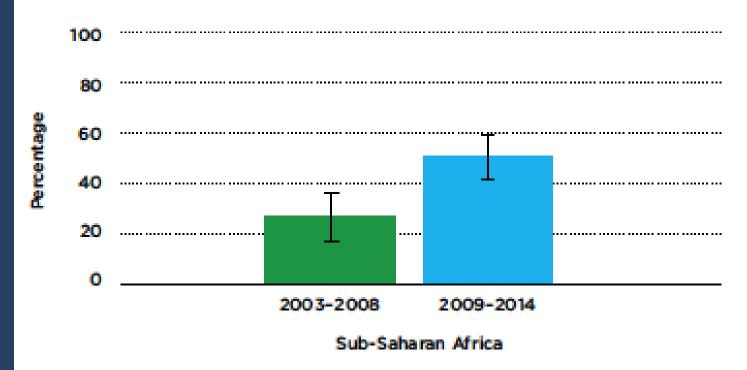
Addressing the barriers

- HIV testing
- Late presentation
- Low ART coverage
- Retention in care
- Retention in ART
- Financing
- Human rights
- Stigma

The first "90"



Awareness of HIV status among people aged 15-49 living with HIV in sub-Saharan Africa



Source: Analysis based on DHS and the South African National HIV Prevalence Surveys.



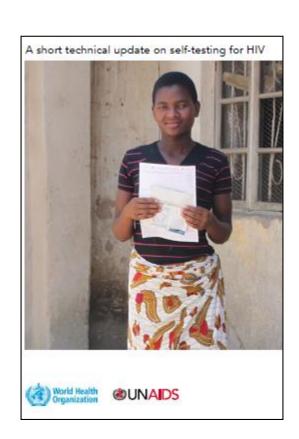
Testing challenges

- Policies and laws
- Stigma and discrimination
- Delivery



A way forward: HIV self-testing

- Available since 1990s
- UNAIDS policy since 2004
- Private non-medical affair
- Convenience and comfort with instant robust results
- Circumvent barriers
- Preferred modality
- heterosexual men, young people, health workers in high prevalence settings, and key populations



A very sensitive issue: identify and treat acute HIV infection, which is responsible for a high proportion of HIV transmission events



The HIV treatment cascade in acutely infected people: informing global guidelines

Sarah E. Rutstein^{a,b,*}, Christopher J. Sellers^{b,*}, Jintanat Ananworanich^{c,d}, and Myron S. Cohen^{b,o}

Purpose of review

Acute and early HIV (AHI) is a pivotal time during HIV infection, yet there remain major shortfalls in diagnosis, linkage to care, and antiretroviral therapy (ART) initiation during AHI. We introduce an AHI-specific cascade, review recent evidence pertaining to the unique challenges of AHI, and discuss strategies for improving individual and public health outcomes.

Recent findings

Presentation during AHI is common. Expanding use of fourth-generation testing and pooled nucleic acid amplification testing has led to improved AHI detection in resource-wealthy settings. Technologies capable of AHI diagnosis are rare in resource-limited settings; further development of point-of-care devices and utilization of targeted screening is needed. Rapid ART initiation during AHI limits reservoir seeding, preserves immunity, and prevents transmission. Reporting of AHI cascade outcomes is limited, but new evidence suggests that impressive rates of diagnosis, linkage to care, rapid ART initiation, and viral suppression can be achieved.

Sum mary

With advancements in AHI diagnostics and strong evidence for the therapeutic and prevention benefits of ART initiated during AHI, improving AHI cascade outcomes is both crucial and feasible. HIV guidelines should recommend diagnostic algorithms capable of detecting AHI and prescribe rapid, universal ART initiation during AHI.

Keywords

acute HIV infection, guidelines, HIV cascade, HIV diagnostics, linkage to care

INTRODUCTION

Acute HIV is a very brief but critical phase of HIV infection. Historically, acute HIV infection has been defined as lasting only until the emergence of HIVspecific antibodies [1], because lack of antibodies in the presence of viral RNA was used as an operational definition of the stage of disease. However, as antibody tests have become ever more sensitive and many patients have been followed from infection onward [2], it is wiser to consider acute and early HIV infection as a package (sometimes referred to as 'primary infection'), because clearly the important events that transpire after infection extend for a longer time than required for antibodies to form [3]. The exact time at which acute and early infection should be considered as 'established' infection has not been resolved. However, the transmission risk associated with acute and early infection lasts at least 4 months [4].

Acute HIV infection and early HIV infection (hereafter referred to as AHI) are associated with extremely high viral loads, seeding of viral reservoirs, and a disproportionate contribution to onward HIV transmission. Failure to diagnose and treat persons with AHI has significant individual and public health implications. Responding to these

"Department of Health Policy and Management, University of North Carolina at Chapel Hil, Chapel Hil, North Carolina, USA, "Division of Infectious Diseases, University of North Carolina at Chapel Hill, Chapel Hill, North Carolina, USA, "U.S. Militany HIV Research Program, Walter Read Amy Institute of Research, Siver Spring, "Henry M. Jackson Foundation for the Advancement of Military Medicine, Bethesda, Maryland and "Department of Epidemiology, University of North Carolina at Chapel Hill, Chapel Hill, North Carolina, USA

Correspondence to Sarah E. Rutstein, PhD, Department of Health Policy and Management, CB #7411, University of North Carolina at Chapel Hill NC 27599-7411, USA. Tel: +1 206 419 8151; fax: +1 919 986 6961: e-mail: sarah nutstein@meduno.edu

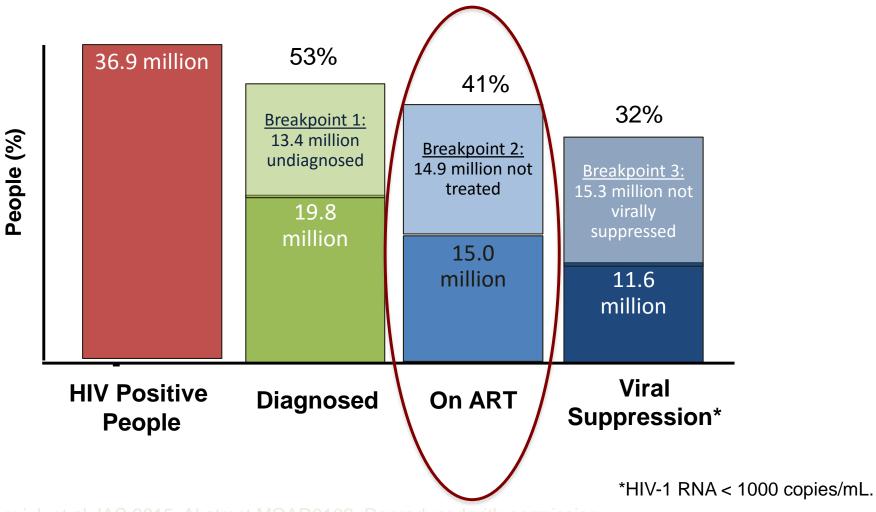
*Sarah E. Rutstein and Christopher J. Sellers contributed equally to the writing of this article.

Curr Opin HIV AID \$ 2015, 10:395-402 DOI:10.1097/COH.0000000000000193

1746-630X Copyright © 2015 Wolters Kluwer Health, Inc. All rights reserved.

www.co-hivandaids.com

The second "90"



Levi J, et al. IAS 2015. Abstract MOAD0102. Reproduced with permission.

National policies regarding initiation of ART (May 2015)

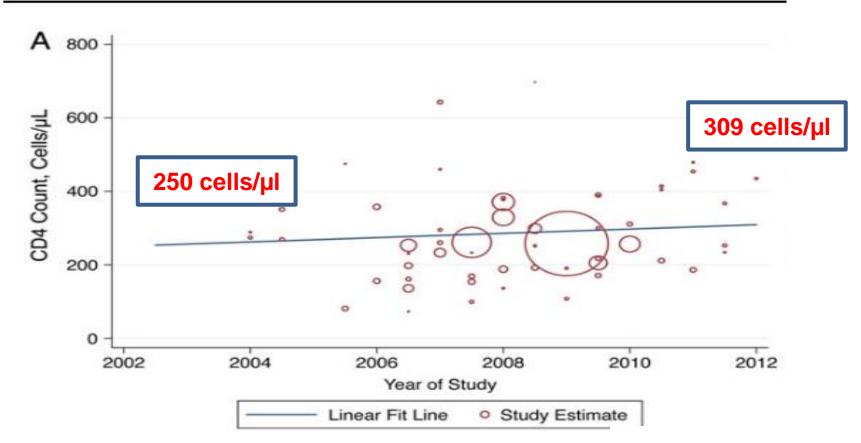


^{*} CD4 cells are key players in the body's immune system that are progressively depleted by HIV infection. Clinicians monitor HIV-related immune deterioration by measuring the number of CD4 cells in a cubic millilitres of plasma.



Late presentation

Trends in CD4 Count at Presentation to Care and Treatment Initiation in Sub-Saharan Africa, 2002–2013: A Meta-analysis



Antiretroviral treatment coverage is still very low in many regions

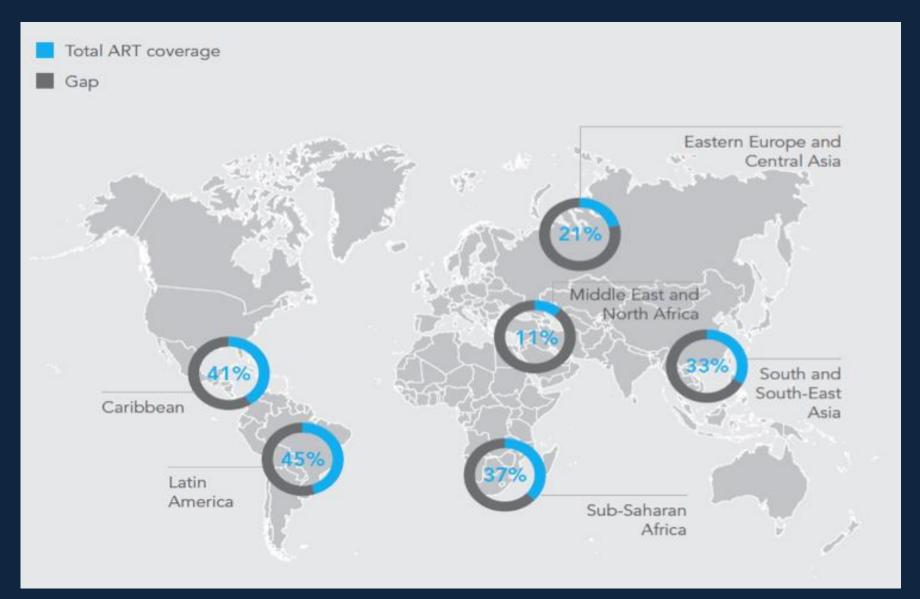
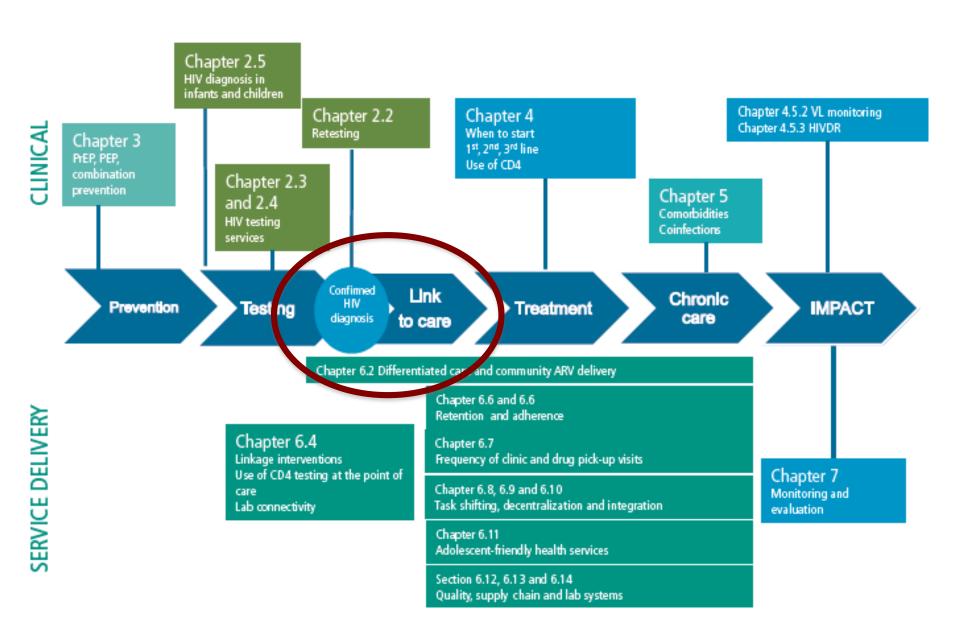


Fig.1. Guidance across the continuum of HIV testing, prevention, treatment and care in the updated Consolidated guidelines on the use of antiretroviral drugs for treating and preventing HIV infection

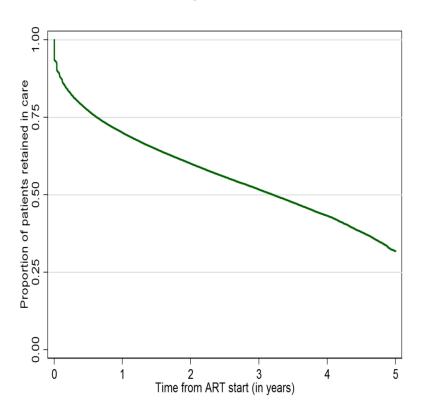


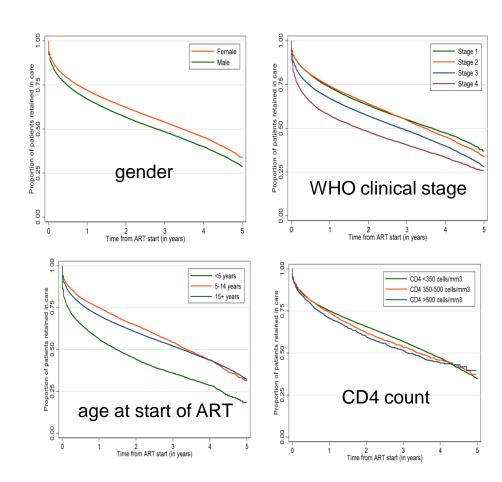
Starting patients on ART is just the first step....

....retaining people in therapy and keep the virus fully suppressed (for years) is far more complicated....

Retention in HIV care programmes

Global analysis of retention in care in initial HIV care and treatment program in the leDEA regions (41 countries)





WHO-leDEA collaboration, 2015



The new ART eligibility criteria will increase the proportion of asymptomatic patients in ART programs. As they are still well, these patients may perceive no short-term benefit from entering treatment, with consequent ART cessation, especially in the face of onerous ART procurement or regimens with persistent side effects.



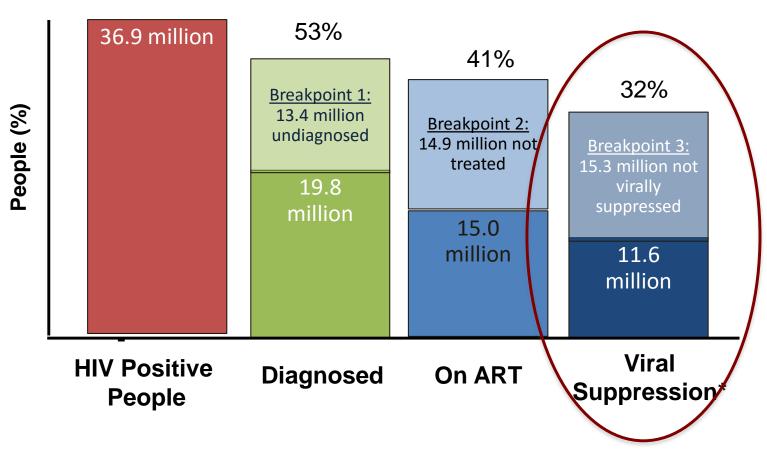
"Why shall I take this pill every day if I am feeling well "?

"Yes, I stopped my medication because I feel better and think I am cured"

Innovative models of HIV care are needed

Simplification of ART delivery, at least for asymptomatic and clinically stable patients, through full community-based care models, including motivational counseling and HIV infection literacy programs run by trained community health workers.

The third "90"



*HIV-1 RNA < 1000 copies/mL.

To retain patients in ART we definitely need innovative models of care but we may also need more tolerable regimens

ART Optimizatio n Strategy	Tolerabilit y	Resistanc e	Conven ience	PW, TB, children	Cost Reduction	What action are needed?	Considered for 2015 WHO guidelines review?
Low dose EFV	✓	?	√	?	√	pK studies (PW & TB)	√
Low dose DRV/r (as FDC)	✓	?	✓	?	✓	 pK studies (titration of best DRV:RTV ratio) RCT (comparative trials: standard vs low dose) 	✓
DTG	✓	✓	✓	?	✓	 Studies in PW, TB & children Comparative trials (TDF /TAF in 1st line) RCT (DRV/r + DTG in 2nd line) 	√
TAF	✓	?	✓	?	✓	 Comparative trials using DTG Studies in PW, TB & children 	*
Long- acting formulation	✓	?	✓	√	WHO Think	Phase II/III studies Tanktreentings թթ√շրկա Optin trials)	nizatio <mark>n (</mark> 2013

WHO goes in the right direction.....

Table 4.1: First-line ART regimens for adults, adolescents, children and pregnant/breastfeeding women

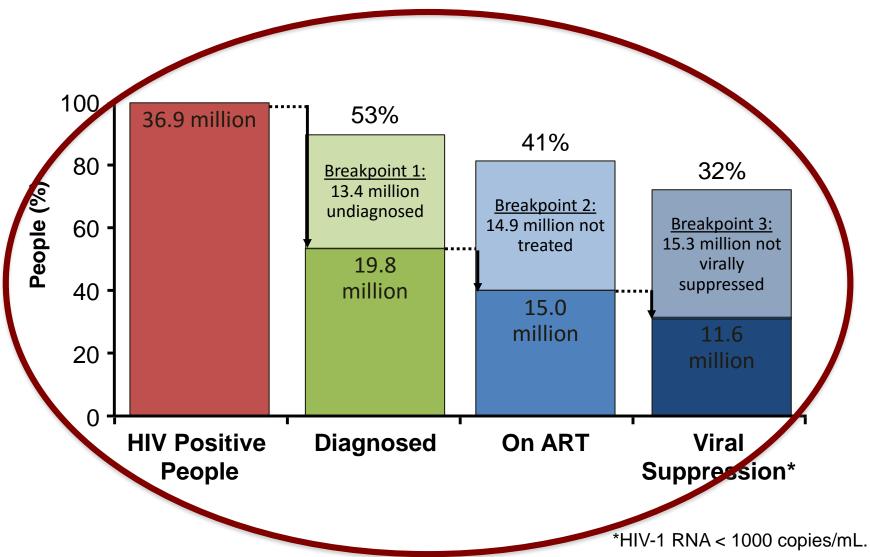
First-line ART	Preferred first-line regimens	Alternative first-line regimens ^{a,b}
Adults and adolescents	TDF + 3TC (or FTC) + EFV	AZT + 3TC + EFV (or NVP) TDF + 3TC (or FTC) + DTG ° TDF + 3TC (or FTC) + EFV ₄₀₀ ° ° TDF + 3TC (or FTC) + NVP
Pregnant/breastfeeding women	TDF + 3TC (or FTC) + EFV	AZT + 3TC + EFV (or NVP) TDF + 3TC (or FTC) + NVP
Children 3 years to less than 10 years	ABC + 3TC + EFV	ABC + 3TC + NVP AZT + 3TC + EFV (or NVP) TDF + 3TC (or FTC) + EFV (or NVP)

ART RETENTION TRIAL

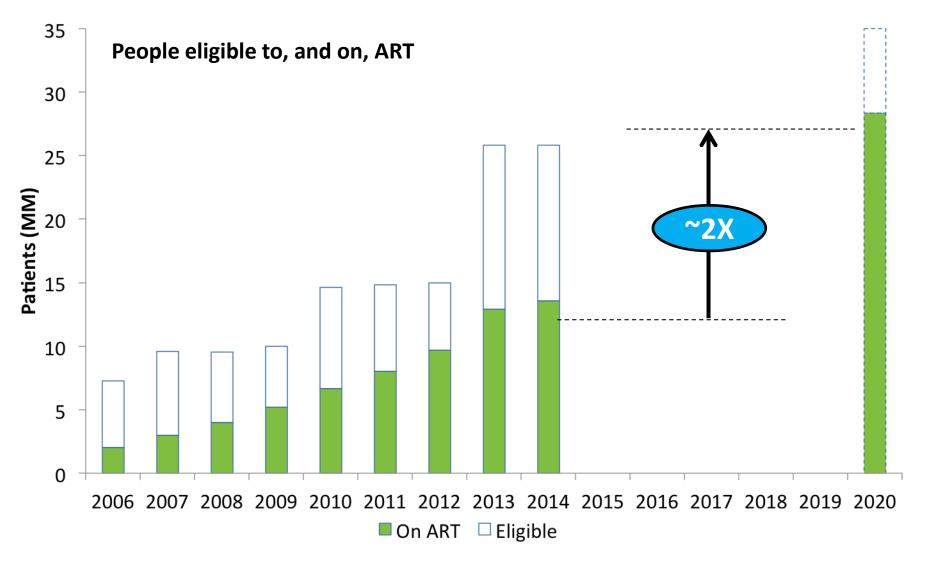
Improving retention in care and adherence to ART of asymptomatic HIV+ patients: a factorial, randomized trial exploring the <u>interaction of two interventions</u>: a community-based HIV-care model combined with a more tolerable first line regimen. (SA, Zimbabwe, Zambia, Rwanda, Ethiopia, UK, Italy)

	Facility-based Care	Community-based Care	Main effect
EFV-based regimen	Interaction effect: 65%	Interaction effect: 75%	EFV-based regimen: 70%
DTG-based regimen	Interaction effect: 75%	Interaction effect: 90%	DTG-based regimen : 82,5%
Main effect	Facility- based care: 70%	Community-based care: 82,5%	

ADDRESSING "OVERARCHING" BARRIES



The next target: almost doubling the number of people on ART



Source: UNAIDS, Global AIDS Report 2006-2013.; WHO UNICEF and UNAIDS, Global Update on HIV Treatment 2013.

FINANCING

THE AIDS RESPONSE STILL REMAINS DONOR DEPENDENT IN MANY COUNTRIES

HIV continues to remain more donor dependent than other health programmes, although HIV expenditure constitutes only a small fraction of total health expenditure across all income groups. Programmes for key populations continue to be mainly funded through international donors, which makes their sustainability questionable once the donors withdraw.

2 SIGNIFICANT FINANCIAL GAPS AT THE COUNTRY LEVEL ARE MADE WORSE BY INEFFICIENT SPENDING

Despite considerable amounts of funding for HIV over the past 15 years, important financial gaps remain in all low- and middle-income countries, with the problem made worse by inefficient allocation and implementation of resources. Across the board, HIV responses remain uneven—variations of unit costs can be observed not only between regions and types of epidemic but also within the same country.

3 FUNDING FOR CIVIL SOCIETY ORGANIZATIONS IS BEING ROLLED BACK

Many civil society organizations are reporting cutbacks in the funding available for core functions such as advocacy, accountability, mobilization, networking and community delivery of services. When current health systems are insufficient for an effective and efficient response, funding of civil society and community organizations is needed more than ever.

4 GLOBALLY, AN ADDITIONAL US\$ 8-12 BILLION NEEDS TO BE AVAILABLE ANNUALLY BY 2020

Increasing funding for treatment is crucial to achieving the goal of ending the AIDS epidemic as a public health threat. Globally, an additional US\$ 8–12 billion needs to be available annually by 2020. Equally important is the need for increased funding for comprehensive programmes for key populations in order to improve access to testing, treatment outcomes, retention in antiretroviral therapy and HIV prevention. Highly efficient use of the resources is a must.

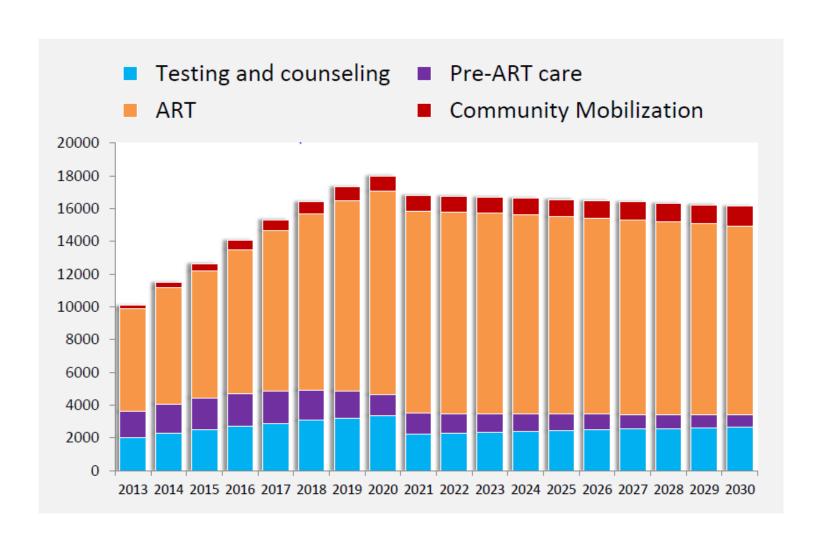
RESOURCES ARE NOT ALWAYS ALLOCATED TO PLACES AND POPULATIONS WHERE THEY WILL MAKE THE MOST IMPACT

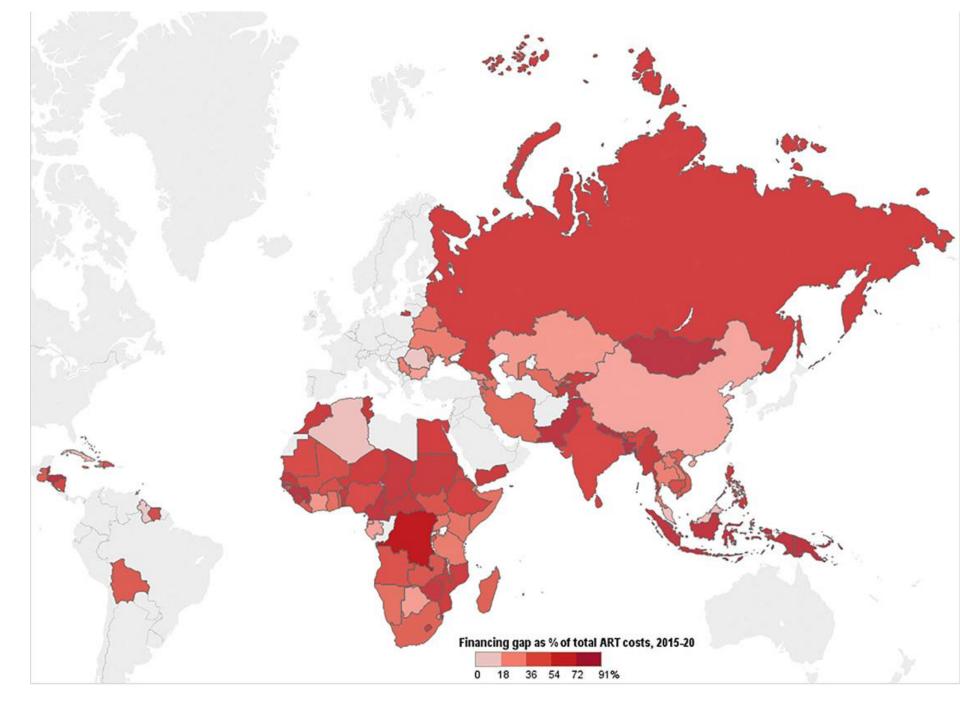
Resources are not always allocated to places and populations where they will make the most impact, and any move towards better allocative efficiency requires careful political negotiations and a full consideration of equity and human rights. The unprecedented funding for the HIV response has generated a vast amount of data, tools, analysis and strategic information about locations and populations, but that does not always translate into policy shifts or changes in how business is done.

Moving towards end of AIDS: main results and new ambitious targets...

Key parameters	2005	2015	2020	2030
New HIV infections	3 million	2 million	500,000	200,000
AIDS-associated deaths	2.4 million	1.2 million	500,000	400,000
PLHIV accessing ART	1.5 million	15 million	30 million	ALL
Investments for global HIV response (US\$)	7 billion	20 billion	32 billion	29 billion

Resource Needs for Treatment, Care and Support US\$ Million





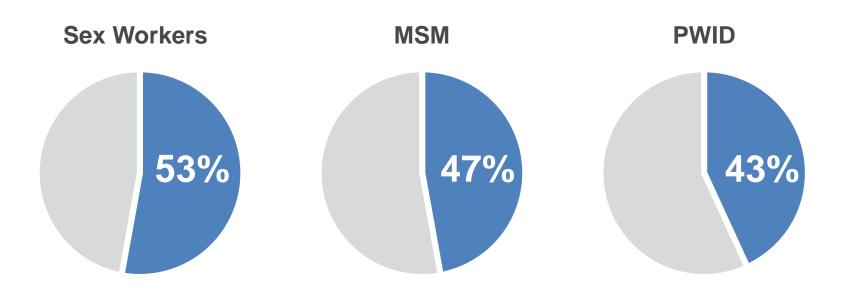
The next barrier: Political committment

- To avoid global investments shifting elsewhere, we definitely need to revitalise community mobilisation and the HIV/AIDS transformative partnership model we used in the past decades.
- We need to make clear to world leaders that we are definitely **not close to the solution**, that without political support to the HIV/AIDS community the window of opportunity which science has opened will be missed, and that the AIDS epidemic is set to grow again **without strong continued financial support** for country programmes.

The next barrier: Human Rights not respected everywhere

- To avoid global investments shifting elsewhere, we definitely need to revitalise community mobilisation and the HIV/AIDS transformative partnership model we used in the past decades.
- We also need to make clear to world leaders that we are definitely **not close to the solution**, that without political support to the HIV/AIDS community the window of opportunity which science has opened will be missed, and that the AIDS epidemic is set to grow again **without strong continued financial support** for country programmes.
- And without human-rights-based decriminalising approaches targeting key populations.

Discriminatory Laws and Policies



Percentage countries whose laws, regulations, or policies can hinder service provision for key populations

obstacles to effective HIV prevention, treatment, care and support services for key populations"

KAP...

- People who use drugs
- Men having sex with men
- Women having sex with women
- Transgender people
- Sex workers
- Incarcerated people
- Displaced people, refugees
- Migrants
- •

Diversity of HIV epidemics: interventions shall be targeted

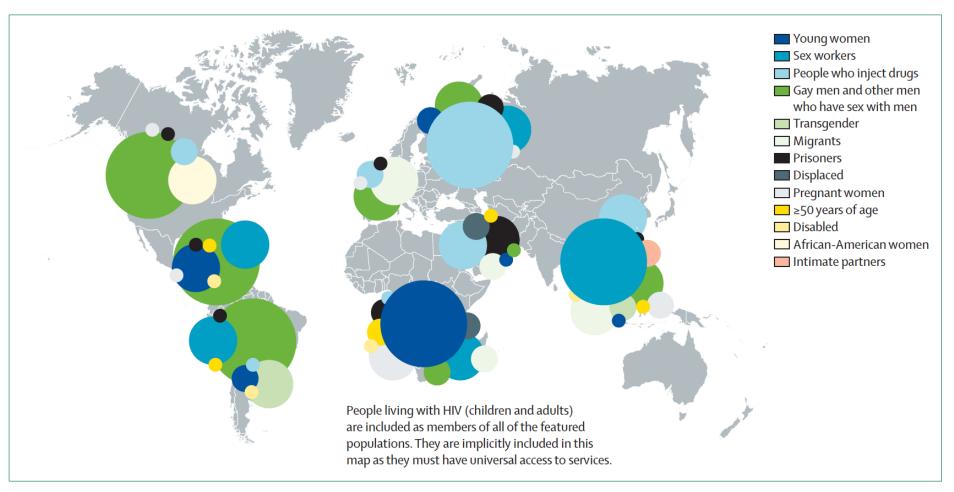


Figure 4: The importance of location and population

Source: The Gap Report.⁴

90/90/90 at a glance

Target component	What it means?	What was achieved until now?	How can it be improved ? (key issues)
The "1st 90"	90% of all PLHIV diagnosed	Approximately 50% of all PLHIV	 Early diagnosis (focus on key populations, adolescents, young men) Innovative testing strategies Reduce stigma//discrimination
The "2 nd 90"	90% of all diagnosed PLHIV on ART (81% of all PLHIV)	Approximately 40% of all PLHIV	 Early ART initiation (ART for all with prioritization) Linkage to care Retention support (care packages) Treatment optimization (new regimens) Reduce stigma//discrimination
The "3 rd 90"	90% of all PLHIV on ART with supressed VL (73% of all PLHIV)	Approximately a 30% of all PLHIV*	 Early detection of treatment failure (expanded access to VL testing) Retention support (adherence/social /community) Treatment optimization (new regimens & maintenance strategies) Reduce stigma//discrimination

^{*} McMahon J. et al." Bulletin of the World Health Organization 91.5 (2013): 377-385.

90 90 90 – Conclusions

- Treatment as Prevention is definitely part of the solution
- The 90 90 90 targets are achievable
- The whole treatment cascade shall be supported.
- Prevention strategies shall be put in place (both behavioral and biomedical)
- Existing barries shall be addressed:
 - Expand testing through innovative strategies
 - Stigma is properly addressed
 - Innovative models of caring HIV as a lifelong disease are implemented
 - Community involvement is extended (and supported)
 - The focusing is on Key Affected Populations
 - Donor support expands
 - Governments tackle stigma, discrimination and protect human rights
 - Research on better medicines, and towards a vaccine and a cure shall continues to make progress

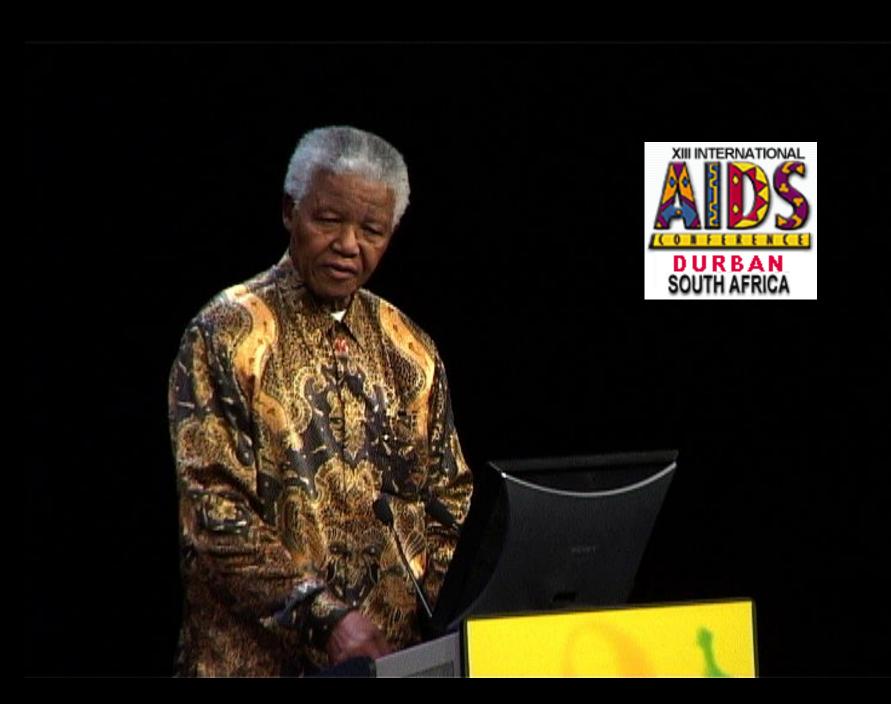
ACHIEVING 90:90:90: A GLOBAL GAME CHANGER FOR PUBLIC HEALTH

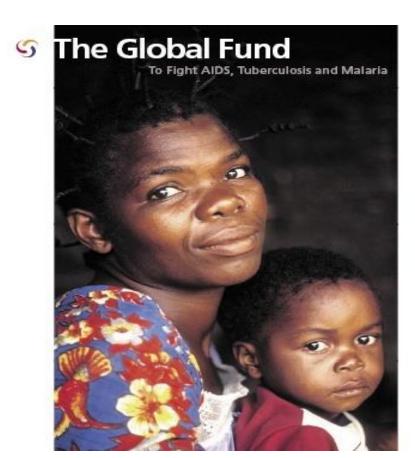
The Goal

The Tools

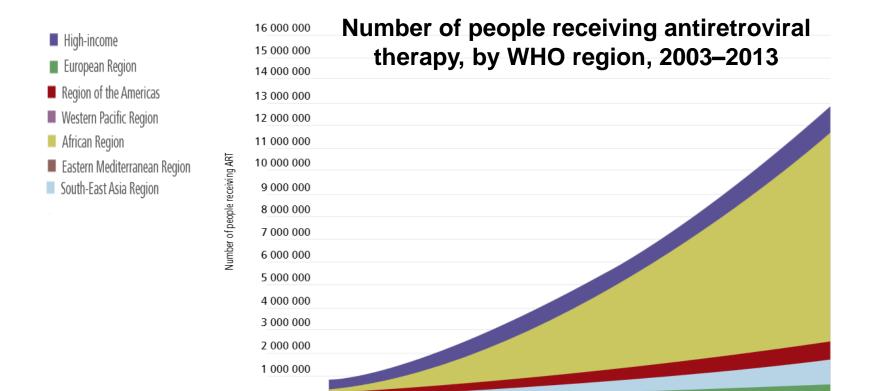
Addressing the barriers

From Durban to Durban









Number of people receiving ART globally rose from ~2 million in 2005 to ~15 million in 2015

Source: Global AIDS Response Progress Report.

From Durban to Durban

From the XIII International AIDS Conference in 2000 to the 2016 Durban conference

From universal access
to treatment and care....
....to ending the HIV epidemic.

From Durban to Durban



Thanks

- Badara Samb
- Anton Pozniak
- Marco Vitoria
- Ethiopian local partners
- The local HIV community
- The operational research group of my Institution