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Tracking viral control in adolescents on antiretroviral therapy in Lusaka, Zambia: A retrospective cohort analysis

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Scan this QR code with your smart phone or mobile device to read online. **Background:** In 2023, an estimated 39.9 million people globally were living with HIV, of which 1.55 million were adolescents aged 10–19 years. The 2021 Zambia HIV Impact Assessment revealed lower viral suppression rates in adolescents (15–24 years old) compared to adults on antiretroviral therapy (ART). Lusaka District, Zambia, has the highest number of adolescents on ART, with a 15.1% HIV prevalence in 2018.

Objectives: To determine the prevalence and factors associated with viral suppression among adolescents living with HIV (10–19 years) on ART in Lusaka District, Zambia.

Method: A retrospective cohort analysis was done of 3409 adolescents on ART at public health facilities in Lusaka from January 2023 to December 2023, and who had viral loads recorded. Socio-demographic, clinical, treatment and behavioural data were extracted from electronic health records and analysed using SPSS version 29.

Results: The adolescent cohort in Lusaka achieved 91.8% viral suppression rate (< 1000 copies/mL), with 79% fully suppressed (< 50 copies/mL). In multivariate analysis, older adolescents (15–19 years) had lower odds of suppression compared to younger adolescents (10–14 years) (adjusted odds ratio [AOR] = 1.79; confidence interval [CI] : 1.32–2.43). Higher odds of viral suppression were linked to first-line dolutegravir regimen (AOR = 5.12; CI: 3.23–8.11) and optimal adherence (AOR = 1.89; CI: 1.03–3.47), while regimen switches reduced the odds of viral suppression (AOR = 0.60; CI: 0.45–0.80).

Conclusion: Zambia reached the previous UNAIDS 90-90-90 targets with a viral suppression rate of 91.8%. However, to reach the revised 95% target by 2030, tailored interventions should be implemented to improve adherence and retention in care, particularly for older adolescents.

Keywords: adolescents; antiretroviral therapy; cohort; viral suppression; retrospective; Zambia.

What this study adds: This study adds insights into adolescent viral suppression rates, highlighting disparities by age and treatment regimen. The study provides evidence that first-line dolutegravir regimens and optimal adherence significantly enhance viral suppression.

Introduction

The HIV pandemic remains a significant global health challenge, particularly affecting adolescents aged 10–19 years. In 2023 an estimated 39.9 million (confidence limits: 36.1–44.6 million) people were living with HIV, of which approximately 1.55 million (1.2–1.9 million) were adolescents.¹ Adolescents and young adults aged 10 years to 24 years continue to be disproportionately affected by HIV with 360 000 (240 000–480 000) new HIV infections, of which 140 000 (39 000–240 000) were older adolescents aged 15 years and 19 years.¹ This population faces a significantly higher risk of antiretroviral therapy (ART) non-adherence, increasing their susceptibility to mortality and morbidity compared to other age groups.²

The Joint United Nations Agency on HIV/AIDS (UNAIDS) 95-95-95 global target, which states that 95% of people living with HIV (PLWH) should know their status, 95% of those who know their status should receive ART, and 95% of those in care should have a suppressed viral load (VL) by 2030 is recommended for low- and middle-income countries as part of a fast-track strategy to end the HIV epidemic.³ Antiretroviral therapy is crucial for improving the duration and quality of life of PLWH by suppressing the virus, reducing morbidity and mortality and enhancing overall health.⁴

The World Health Organization (WHO) recommends that all PLWH on ART undergo routine VL testing to achieve these global targets. At local level, VL monitoring is critical for measuring ART efficacy, early detection of drug resistance, improving clinical outcomes, improving treatment adherence and reducing HIV transmission.⁵ Zambia has adopted the WHO recommendation of regular monitoring of VL, with the first VL test at 6 months and the second 12 months after starting ART, and annual testing thereafter.⁵

The WHO recommends three key categories for HIV VL assessments: unsuppressed (> 1000 copies/mL); suppressed (< 1000 copies/mL); and fully suppressed or undetectable (VL not detected by test or sample type used) < 50 copies/mL.⁶ Zambia adopted the WHO guidelines that set viral suppression as not suppressed at \geq 1000 copies/mL and suppressed as < 1000 copies/mL.^{7,8} However, while Zambia has set viral suppression at < 1000 copies/mL, it is important to note that maintaining VL levels below 200 copies/mL is critical to effectively preventing HIV transmission, aligning with the 'Undetectable = Untransmittable (U = U)' principle. This is particularly relevant among adolescents, where ensuring sustained viral suppression at this lower threshold can significantly reduce transmission risks and improve long-term health outcomes.⁹

Adolescents living with HIV (ALHIV) face unique challenges that are associated with rapid physical, psychological and physiological changes during adolescence that impact behaviour negatively on treatment adherence,^{10,11} which in turn leads to mental health problems,^{12,13} reduced engagement with healthcare and lower rates of viral suppression. However, the adolescent population is often overlooked because programmatic reporting does not use the WHO definition of adolescents (10–19 years old) and instead classifies and reports them under paediatrics (0–14 years old) or adult (15 years old and older) populations.

Viral suppression in PLWH is influenced by a complex interplay of socio-demographic, clinical, treatment and behavioural factors. Existing literature emphasises the role of socio-demographic characteristics, such as (current) age, age at ART initiation and gender in determining viral suppression outcomes.^{14,15} Several studies report lower rates of viral suppression in older adolescents compared to younger adolescents due to treatment fatigue (being on ART from childhood), and not being adequately prepared to transition to adult care and self-management.^{16,17} Similar conflicting results are reported for female versus male adolescents due to gender differences in health-seeking behaviour,¹⁸ as well as higher infection rates amongst adolescent girls compared to boys.¹

Clinical factors such as duration on ART, and baseline CD4 count and WHO clinical stage have been found to be strong predictors of viral suppression.^{14,17,19,20} Studies have shown that adolescents who start ART with a less advanced immunodeficiency (CD4 > 200 cells/mm³ – 350 cells/mm³)

appear to have better virological outcomes than adolescents who start with a more severe immunodeficiency (CD4 count < 200 cells/mm³).^{19,21,22,23,24} However, other studies found that higher CD4 counts were not associated with viral suppression.^{25,26,27} Various studies have reported variations around the duration on ART with viral suppression, where longer duration on ART is expected to be associated with viral suppression.²⁸

Treatment characteristics such as the choice of antiretroviral regimen and changes to a more efficacious regimen are vital for achieving viral suppression.^{29,30,31} Following the WHO recommendion for use of dolutegravir (DTG) as the preferred core agent in ART for first and second-line regimen,³² studies have reported improved viral suppression in PLWH which includes children and adolescents.³³

Behavioural characteristics, such as optimal adherence to medication and retention in care, are strongly correlated with viral suppression.³⁴ Optimal adherence is defined as taking at least 95% medications at prescribed times and frequencies (at least 95%),^{35,36} while retention in care is a patient's regular engagement with medical care at a health care facility after initial entry into HIV clinical care as set out with appointments schedules.³⁷ According to Zambian guidelines, one is deemed out of care (not retained) once they miss a pharmacy refill for 30 days or more.⁷

Whereas the risk factors influencing viral suppression among ALHIV are known and clearly illustrated in the literature, these factors may differ in various contexts. It is therefore essential to identify the specific set of factors associated with viral suppression that are prevalent in ALHIV in Zambia, as this information is crucial for tailoring interventions and programmes to improve adherence and viral suppression as progressive steps to reach the UNAIDS 95-95-95 targets by 2030.

Objectives

The objective of the current study was to determine the prevalence and factors associated with viral suppression among ALHIV (10–19 years) on ART from January 2023 to December 2023 in Lusaka District, Zambia.

Research methods and design Study design

We conducted a retrospective study of routinely collected data from ALHIV on ART from January 2023 to December 2023 to determine the prevalence and factors associated with viral suppression in the Lusaka District. Routine sociodemographic, clinical, treatment, and behavioural data were collected for adolescents aged 10 years to 19 years who were enrolled in public health facilities providing ART in Lusaka District. Behavioural data included measures of optimal adherence (assessed through pharmacy refill data) and retention in care (assessed through missed appointment tracking).

Study context

Lusaka District is located in the southern part of Zambia's Central Province and serves as the capital city of the country. Health services in the district are provided at different levels of care. The levels of care in the public sector range from health posts (lowest level) to specialised hospitals which are designated third-level hospitals (highest level). The master facility list shows that Lusaka District has 36 health posts, 31 health centres, five level one hospitals and eight level three (specialised) hospitals. Health posts and health centres mainly provide primary healthcare services which include HIV services. First level hospitals act as referral facilities for health posts and health centres and also provide HIV services. Third level hospitals are specialised facilities which mainly handle complicated cases referred to from lower levels of care.

Study population and sampling

The study population consisted of all adolescents aged 10–19 years receiving ART and accessing ART services in Lusaka District in 2023.

Data collection

Routine data were extracted from an electronic health record (EHR) called SmartCare. SmartCare is a commonly used EHR in public health facilities across Zambia. Its implementation varies by facility: some operate on an electronic last or e-last model, where data entry is completed retrospectively by designated data entry personnel, while others use an 'electronic first' (e-first) approach, where healthcare providers at all service points within the facility input patient data in real time.

Study data were extracted in August 2024 and consisted of anonymised data identified by unique patient identifiers. Data included demographic, clinical, treatment, and behavioural data collected during routine clinical visits for patient management and monitoring of all patients receiving ART. All ART data are captured using SmartCare and therefore any missing data were unable to be retrieved. Viral load was the primary outcome variable: VL < 50 RNA copies/mL was categorised as 'fully suppressed'; VL > 50 copies/mL – 999 copies/mL was categorised as suppressed; and VL > 1000 copies/mL was categorised as 'unsuppressed' according to national clinical ART guidelines.⁷ In this study, 'viral suppression' applies to all participants who had a VL of < 1000 copies/mL most recently recorded in SmartCare.

Socio-demographic variables included gender, current age, and age at ART initiation. Clinical characteristics included baseline WHO clinical stage, CD4 count (baseline and current), pregnancy, lactation, and history of active tuberculosis (TB). Treatment characteristics included current ART regimen, current ARV regimen class, change in ART regimen, and use of TB preventive therapy. Behavioural characteristics included retention in care and optimal adherence to ART. In the current study, retention in care was defined as currently being active on ART, and optimal adherence to ART is defined as missing a medication refill appointment by no more than 2 days.

Data analysis

The extracted data were entered into an Excel spreadsheet, cleaned, and then imported into the IBM Statistical Package for the Social Sciences (SPSS) for Mac, Version 29.0.2.0 (IBM Corp., Armonk, New York). For inferential analysis, variables with missing values were treated using complete-case analysis, with cases with missing data excluded from the analysis. For categorical variables, descriptive frequency tables were created to describe the socio-demographic, clinical, behavioural, and treatment characteristics of the study population. Bivariate analysis was used to determine the significance and strength of associations between the independent variables and the outcome variable (VL < 1000 copies/mL). Chi-square tests and crude odds ratios were used to examine and quantify associations, with significance set at P < 0.05. Variables that showed significant associations with viral suppression during bivariate analysis, at a significance level of 5%, were included in the multivariate analysis to identify factors independently associated with viral suppression, and presented as adjusted odds ratio (AOR) with 95% confidence intervals (95% CI).

Ethical considerations

Ethical clearance was obtained from the University of the Western Cape Biomedical Research Ethics Committee (reference number: BM24/3/4), and the Mulungushi University School of Medicine Ethics Committee (reference number: SMHS-MU2-2024-04) and the Zambia National Health Research Authority (reference number: NHRA1186/15/05/2024). Approval for the study and access to the data were obtained from the Zambia Ministry of Health (reference number: MH/101/22/3). During data extraction, the adolescents' unique identities, such as ART number, identity number, first and last name, were excluded to ensure complete anonymity and protection of personal information.

Results

Realisation of sample

Records of 128 991 PLWH attending ART services in health facilities in Lusaka District from January 2023 to December 2023 were obtained from the EHR. Figure 1 shows that of 128 991 records, 125 013 were records of PLWH aged less than 10 years and more than 19 years who were in care in 2023, while 3978 records were of adolescents aged 10 years to 19 years during 2023. 569 records of the 3978 adolescents had missing VL results in the EHR (see Figure 2).



ART, antiretroviral therapy; EHR, electronic health record; VL, viral load. **FIGURE 1:** Flow chart of participants selected in the study.

Socio-demographic, clinical, treatment and behavioural characteristics

Table 1 shows the socio-demographic, clinical, treatment and behavioural characteristics of ALHIV enrolled in the ART programme in public health facilities in Lusaka District, Zambia.

This study involved 3409 adolescents (aged 10 years to 19 years) with a mean age of 14.97 years (standard deviation [s.d.] = 1.15); of these, 53% (1820) were female. The mean age at ART initiation was 7.51 years (s.d. = 1.15). The majority of participants (2935, 98.7%) were at WHO clinical stage 1. Most participants (1897, 78% and 1897, 88.6%) had both baseline and recent CD4 counts \geq 350 cells/mm³. The vast majority of participants (3241, 95.1%) received first-line therapy, and were on a DTG-based regimen (3351, 98%). Majority of participants (3132, 91.9%) remained in care, while 9.1% (277) experienced an interruption in treatment (IIT), transferred out or died during the study period. Most participants (2837, 96.5%) demonstrated optimal adherence to their ART, as defined by not missing a pharmacy refill by more than 2 days.

Determinants of viral suppression

The overall viral suppression was 91.8% (< 1000 copies/mL) while fully suppressed was 66% (< 50 copies/mL). Table 2 shows that in bivariate analysis, the odds of viral suppression were significantly higher (94% vs 90%; P < 0.01) in individuals aged 10 years to 14 years than those aged 15 to 19 years (crude OR = 1.79, 95% CI = 1.32 – 2.42). After adjusting for other variables, the significant association between viral suppression and age remained (adjusted OR [AOR] = 1.79, 95% CI = 1.32 – 2.43) (see Table 2). No statistically significant associations were found between viral suppression and gender (P = 0.539), and age at initiation of ART (P = 0.2) (see Table 1).

There were statistically significant associations between viral suppression and pregnancy (P = 0.010), baseline CD4 count (P = 0.011) and latest CD4 count (P < 0.01). However, CD4 counts were only recorded for 42% and 54% of



FIGURE 2: Distribution of viral load records and suppression status.

participants at baseline and current (observation year), respectively, while pregnancy and breastfeeding were documented for only 3% and 53% of all female participants and were therefore excluded from the multivariate analysis. No statistically significant associations were identified between viral suppression and clinical factors such as history of active TB (P = 0.145), current WHO stage (P = 0.486), and breastfeeding (P = 0.559) (see Table 1).

Statistically significant associations between viral suppression and treatment factors, including current ART regimen (P < 0.01), current ARV regimen class (P < 0.01) and change in ART regimen (P < 0.01) were observed. In the multivariate logistic regression, the relationship between viral suppression and current ART regimen was statistically significant (AOR = 6.14, 95% CI = 3.86 – 9.48), while the relationship between viral suppression and current ART regimen class did not retain statistical significance. In multivariate logistic regression, the relationship between viral suppression and change in ART regimen retained statistical significance (AOR = 0.62, 95% CI = 0.46 – 0.83).

The relationship between viral suppression and behavioural characteristics optimal adherence (AOR = 1.82, 95% CI = 0.97 - 3.41) and retention in care (AOR = 1.25, 95% CI = 0.93 - 1.69) did not retain statistical significance after adjusting for potential confounders.

Discussion

Lusaka District made significant progress towards achieving the previous goal of 90-90-90 set by UNAIDS, reaching a 91.8% viral suppression rate for ALHIV, based on a threshold of < 1000 copies/mL. Similar findings were reported from Tanzania³⁸ and South Africa³⁹ with 89.9% and 88%, respectively. These consistent results across diverse settings underscore the effectiveness of targeted ART programmes for adolescents as propagated by UNAIDS.⁴⁰ However, to further reduce transmission among this age group, adopting the more rigorous viral suppression threshold of < 50 copies/ mL, referred to as 'fully suppressed' and recommended by the WHO, would be a critical consideration. TABLE 1: Determinants of viral suppression (< 1000 copies/mL) among adolescents on antiretroviral therapy in the Lusaka District, 2023 (N = 3409).

Characteristics	Total		Viral suppression				Р
			Y	'es	N	0	_
	n	%	п	%	n	%	_
Total	3409	-	3132	91.8	277	9.2	-
Age (years)			-	-	-	-	< 0.010*
10–14	1384	40.6	1303	94.1	81	5.9	-
15–19	2025	59.4	1829	90.3	196	9.7	-
Gender	-	-	-	-	-	-	0.539
Male	1589	46.6	1455	91.6	134	8.4	-
Female	1820	53.4	1677	92.1	143	7.9	-
Age at ART initiation (vears)	-	-	-	-	-	-	0.200
0-4	1061	31.1	984	92.4	77	7.3	-
5-9	1178	34.6	1084	92.0	94	8	-
10–14	933	27.3	854	91.5	79	85	-
15-19	237	7.0	210	88.6	27	11 /	
Duration on ART (months)	257		-	-	-	-	0 534
6_12	8	0.2	7	87.5	1	12 5	0.554
12_24	112	2.2	, 100	80.2	12	10.7	_
> 24	2200	06.5	2025	02.0	264	20.7	
Current APT regimen	5269	90.5	5025	92.0	204	0.0	- 0.010*
Eirst line	-	-	-	-	-	-	< 0.010
Second line	146	95.1	01	53.4	214	0.0	-
Second-line	140	4.3	91	62.3	55	37.7	-
	22	0.0	14	03.0	٥	30.4	-
Current ARV regimen class	-	-	-	-	-	-	< 0.010*
	6	0.2	4	66.7	2	33.3	-
DTG-based	3351	98.3	3102	92.6	249	7.4	-
PI-based	30	1.0	12	40.0	18	60.0	-
Unclassified (Incomplete regimen)	22	0.5	14	63.6	8	36.4	-
Change in ART regimen	-	-	-	-	-	-	< 0.010*
Yes	1030	30.2	912	88.5	118	11.5	-
No	2379	69.8	2220	93.3	159	6.7	-
Pregnant† (N = 1820)	-	-	-	-	-	-	0.010*
Yes	17	1.0	13	76.5	4	23.5	-
No	30	1.6	25	83.3	5	16.7	-
Unknown	1773	97.4	1639	92.4	134	7.6	-
Breastfeeding (N = 1820)	-	-	-	-	-	-	0.559
Yes	4	0.2	4	100.0	0	0.0	-
No	1816	99.8	1673	92.1	143	7.9	-
Baseline CD4 count (cells/mm ³)† (N = 1671)	-	-	-	-	-	-	0.011*
< 200	150	9.0	129	86.0	21	14.0	-
200–349	217	13.0	197	90.8	20	9.2	-
≥ 350	1304	78.0	1211	92.9	93	7.1	-
Latest CD4 count (cells/mm³)† (N = 2139)		-	-	-		-	< 0.010*
< 200	74	3.4	60	81.1	14	18.9	-
200–349	168	7.9	147	87.5	21	12.5	-
≥ 350	1897	88.7	1758	92.7	139	7.3	-
Current WHO stage ($N = 2971$)	-	-	-	-	-	-	0.486
Stage I	2935	98.7	2701	92.0	234	8.0	-
Stage II	21	1.0	18	85.7	3	14 3	-
Stage III	14	0.3	14	100.0	0	0.0	-
Stage IV	1	0.0	1	100.0	0	0.0	-
Received TB prophylaxis		-	-	-	-	-	0.053
Ves	2747	80.6	2536	92.3	211	77	-
No	662	19.4	596	90.0	66	10.0	-
History of active TB	-	17.7	550		-	-	0 536
Ves	159	4.7	144	90.6	15	9.4	0.550
No	3250	4.7	2022	91.0	262	9.4 8.1	
Retention in care	5250	33.5	2300	51.5	202	0.1	0.010*
	-	- 01 0	- 1117	-	2020	-	0.015
	2122	91.9	70	55.4	2020	91.1 91.1	-
Ontimal adherance (M = 2040)	2//	9.1	79	0.0	199	6.9	-
Vec	-	-	-	-	-	-	0.044 ~
No	2037	90.5	2009	92.0	228	0.U 12.C	-
NU	103	5.5	09	00.4	14	13.0	-

ART, antiretroviral therapy; ARV, antiretroviral drugs; CD4, cluster of differentiation 4; DTG, dolutegravir; IIT, Interruption in treatment; NNRTI, non-nucleoside reverse transcriptase inhibitor; PI, protease inhibitor; TB, tuberculosis; TO, transfer out. †, Variables not included in the regression model due to very small sample sizes.

*, Statistically significant at p < 0.05

TABLE 2: Determinants of viral suppression of among adolescents (10–19 years) on antiretroviral therapy in Lusaka District, 2023.

Characteristics	Total	Crude OR	95% CI	Adjusted OR	95% CI
Age (years)					
10–14	1384	Ref	-	Ref	-
15–19	2025	1.74	1.32-2.26*	1.69	1.25-2.29*
Current ART regimen					
First-line	3241	Ref	-	Ref	-
Second-line	146	8.55	5.9-12.28*	6.14	3.86-9.48*
Current ARV regimen class					
NNRTI	6	Ref	-	Ref	-
DTG-based	3351	0.16	0.29-0.88*	0.39	0.56-2.74
PI-based	30	-	-	1.27	0.16-10.10
Optimal adherence					
Yes	2837	Ref	-	Ref	-
No	103	1.80	1.01-3.21*	1.82	0.97-3.41
Retention in care					
Yes	3132	Ref	-	Ref	-
No	277	1.38	1.05-1.81	1.25	0.93-1.69

ARV, antiretroviral drugs; CI, confidence interval; OR, odds ratio; DTG, dolutegravir; NNRTI, non-nucleoside reverse transcriptase inhibitor; PI, protease inhibitor. *. Statistically significant at p < 0.05

*, statistically significant at p < 0.05

Of concern in Lusaka is that 15% of adolescents on ART in our study did not have VL results recorded, potentially underestimating or overestimating the true viral suppression levels within the population. This missing VL data may stem from delays in sample transport, missed data entries, or challenges with patient follow-up and adherence to monitoring schedules. Such issues have significant health system implications, including delayed interventions, undermined data-driven decision-making, weakened public health surveillance, misallocation of resources, and fragmented care.

Additionally, our study revealed that older adolescents aged 15–19 years were at greater risk of viral non-suppression compared to those aged 10–14 years, a finding consistent with other studies that link this disparity to challenges in transitioning from paediatric to adult care, which can disrupt continuity of care, reduce adult or caregiver support, and exacerbate adherence issues. Furthermore, adolescence is a period of significant physical, emotional, and social changes that can negatively impact treatment adherence, compounded by treatment fatigue among older adolescents who have been on therapy since early childhood, making it increasingly difficult to sustain adherence over time.^{41,42} Similar findings were reported in South African studies in the Cape Town Metro in the Western Cape Province and Ehlanzeni District in Mpumalanga.^{14,43}

The majority of adolescents currently receiving ART in this study were on first-line treatment (DTG-based) and were more than five times more likely to have viral suppression than those receiving second-line treatment. The high viral suppression rate observed in the current study may be related to the successful introduction of a DTG-based regimen as part of the first-line regimen for all PLWH. This finding is supported by several other studies that reported that the DTG-based regimen is more effective in suppressing VL than the non-DTG-based regimen.^{44,45,46}

Our study found that an optimal adherence rate (of > 95%) was significantly associated with viral suppression. Adherence may have been higher because pharmacy refills based on the EMR were used to assess adherence. However, research suggests that in resource-limited settings, pharmacy refill records correlate well with HIV outcomes and provide a more accurate measure of adherence than patient self-report and clinic-based pill counts.⁴⁷

Limitations

This study is a retrospective cohort analysis based on patient data routinely collected during clinic visits and interactions. As a result, the accuracy and completeness of the data available for extraction depended largely on the robustness of data collection and the thoroughness with which health workers and data entry clerks recorded individual client information. Certain variables, such as CD4 count, pregnancy status, and breastfeeding, had high levels of missing data, resulting in insufficient entries to reliably include them as covariates in the analyses. This incompleteness impacted the study's ability to fully assess the influence of these variables on patient outcomes.

Despite possible limitations, such as missing or incomplete data, routinely collected and tracked data provide a realtime view of service delivery and patient outcomes, supporting timely decision-making and improvements in care. This approach is cost-effective, reduces the need for additional data collection and reflects the everyday reality of healthcare. It enables trend assessment, progress tracking, and identification of emerging issues. Finally, both operational strengths and weaknesses are highlighted while assessing the impact of interventions and system gaps.⁴⁸

Conclusion

Viral suppression among adolescents on ART was relatively high in this study in comparison to some regions, but it was still below the new target of 95% set by the UNAIDS. The reported low retention in care rates found in our study warrant further investigation and possible intervention. Older adolescents (15–19 years), those on second-line regimens, and non-adherent adolescents were found to be at higher risk of viral non-suppression. The results highlighted the need for targeted interventions focused on adolescentfriendly services which may include age-appropriate health services, psychosocial support which includes mental health services and disclosure and intentional transition processes from paediatric to adult care.

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Competing interests

The authors declare that they have no financial or personal relationships that could have inappropriately influenced them in writing this article.

Authors' contributions

K.M., B.v.W. and T.C. contributed to the design and implementation of the research; K.M. conducted the statistical analysis under supervision of B.v.W. and T.C. and wrote the first draft. K.M., B.v.W. and T.C. contributed to subsequent drafts and approved the final article.

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Data availability

The data that support the findings of this study are available from the corresponding author, K.M., upon reasonable request.

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References

- UNICEF. Adolescent HIV prevention [homepage on the Internet]. 2024 [cited 2024 Jul 05]. Available from: https://data.unicef.org/topic/hivaids/ adolescents-young-people/
- UNICEF. Increased screening for adolescents at high risk of antiretroviral non-adherence [homepage on the Internet]. Accel Hub; 2017 [cited 2024 Aug 15]. Available from: https://www.unicef.org/esa/media/10691/file/AH-HEADSS-UNIC EF-Policy-Brief-2022.pdf
- UNAIDS. Understanding fast-track: Accelerating action to end the AIDS epidemic by 2030 [homepage on the Internet]. 2016 [cited 2024 Aug 02]. Available from: https://www.unaids.org/sites/default/files/media_asset/201506_JC2743_ Understanding_FastTrack_en.pdf
- Zurbachew Y, Hiko D, Bacha G, Merga H. Adolescent's and youth's adherence to antiretroviral therapy for better treatment outcome and its determinants: Multicenter study in public health facilities. AIDS Res Ther. 2023;20(1):4–9. https://doi. org/10.1186/s12981-023-00588-y
- WHO. Consolidated guidelines on the use of antiretroviral drugs for treating and preventing HIV Infection [homepage on the Internet]. 2016 [cited 2024 Aug 24]. Available from: https://www.who.int/publications/i/item/9789241549684
- WHO. The role of HIV viral suppression in improving individual health and reducing transmission: Policy brief [homepage on the Internet]. Geneva; 2023 [cited 2024 Aug 10]. Available from: https://www.who.int/publications/i/item/9789240055179
- MOH. Zambia consolidated guidelines for treatment and prevention of HIV infection [homepage on the Internet]. 2022 [cited 2024 Sep 01]. Available from: https://www.differentiatedservicedelivery.org/wp-content/uploads/August-2022-Zambia-Consolidated-Guidelines.pdf
- MOH/National AIDS & STI Control Program. Kenya HIV prevention and treatment guidelines [homepage on the Internet]. 2022 [cited 2024 Sep 01]. Available from: https://www.differentiatedservicedelivery.org/wp-content/ uploads/Kenya-ARV-Guidelines-2022-Final-1.pdf

- Cairns G. The amber light: The World Health Organization's position when an HIV viral load is 'suppressed but not undetectable' | aidsmap [homepage on the Internet]. aidsmap; 2023 [cited 2025 Jan 21]. Available from: https://www. aidsmap.com/news/jul-2023/amber-light-world-health-organizations-positionwhen-hiv-viral-load-suppressed-not
- Ferrand RA, Briggs D, Ferguson J, et al. Viral suppression in adolescents on antiretroviral treatment: Review of the literature and critical appraisal of methodological challenges. Trop Med Int Health. 2016;21(3):325–333. https:// doi.org/10.1111/tmi.12656
- 11. Steinberg L. A social neuroscience perspective on adolescent risk-taking. Dev Rev. 2008;28(1):78. https://doi.org/10.1016/j.dr.2007.08.002
- Spear LP. The adolescent brain and age-related behavioral manifestations. Neurosci Biobehav Rev. 2000;24(4):417–463. https://doi.org/10.1016/S0149-7634(00)00014-2
- Lowenthal ED, Bakeera-Kitaka S, Marukutira T, Chapman J, Goldrath K, Ferrand RA. Perinatally acquired HIV infection in adolescents from sub-Saharan Africa: A review of emerging challenges. Lancet Infect Dis. 2014;14(7):627–639. https:// doi.org/10.1016/S1473-3099(13)70363-3
- Okonji EF, Van Wyk B, Mukumbang FC, Hughes GD. Determinants of viral suppression among adolescents on antiretroviral treatment in Ehlanzeni district, South Africa: A cross-sectional analysis. AIDS Res Ther. 2021;18(1):1–9. https:// doi.org/10.1186/s12981-021-00391-7
- Ntombela NP, Kharsany ABM, Soogun A, et al. Viral suppression among pregnant adolescents and women living with HIV in rural KwaZulu-Natal, South Africa: A cross sectional study to assess progress towards UNAIDS indicators and Implications for HIV Epidemic Control. Reprod Health. 2022;19(1):1–13. https:// doi.org/10.1186/s12978-022-01419-5
- Wyk B Van, Kriel E, Mukumbang F. Retention in care for adolescents who were newly initiated on antiretroviral therapy in the Cape Metropole in South Africa. South Afr J HIV Med. 2020;21(1):1–8.
- Mwangi A, Van Wyk B. Factors associated with viral suppression among adolescents on antiretroviral therapy in Homa Bay County, Kenya: A retrospective cross-sectional study. HIV AIDS (Auckl). 2021;13:111–1118. https://doi. org/10.2147/HIV.S345731
- Mabizela S, Van Wyk B. Viral suppression among adolescents on HIV treatment in the Sedibeng District, Gauteng province. Curationis. 2022;45(1):8. https://doi. org/10.4102/curationis.v45i1.2312
- Elashi BAY, Van Wyk BE. Factors associated with viral suppression among adolescents on antiretroviral therapy in Free State province, South Africa. South Afr J HIV Med. 2022;23(1):5–9. https://doi.org/10.4102/sajhivmed.v23i1.1356
- Maena J, Banke-Thomas A, Mukiza N, et al. Determinants of viral load nonsuppression among adolescents in Mbale District, Eastern Rural Uganda. AIDS Res Ther. 2021;18(1):91. https://doi.org/10.1186/s12981-021-00408-1
- Bennett KK, DeGruttola VG, Marschner IC, Havlir DV, Richman DD. Baseline predictors of CD4 T-lymphocyte recovery with combination antiretroviral therapy. J Acquir Immune Defic Syndr. 2002;31(1):20–26. https://doi.org/10.1097/ 00126334-200209010-00003
- Bonnet F, Thiébaut R, Chêne G, et al. Determinants of clinical progression in antiretroviral-naive HIV-infected patients starting highly active antiretroviral therapy. Aquitaine Cohort, France, 1996-2002. HIV Med. 2005;6(3):198–205 https://doi.org/10.1111/j.1468-1293.2005.00290.x
- 23. Desta AA, Woldearegay TW, Futwi N, et al. HIV virological non-suppression and factors associated with non-suppression among adolescents and adults on antiretroviral therapy in northern Ethiopia: A retrospective study. BMC Infect Dis. 2020;20(1):1–10. https://doi.org/10.1186/s12879-019-4732-6
- Grabar S, Moing V Le, Goujard C, et al. Response to highly active antiretroviral therapy at 6 months and long-term disease progression in HIV-1 infection. J Acquir Immune Defic Syndr. 2005;39(3):284–292. https://doi.org/10.1097/01. qai.0000160925.33935.72
- Bayu B, Tariku A, Bulti AB, Habitu YA, Derso T, Teshome DF. Determinants of virological failure among patients on highly active antiretroviral therapy in University of Gondar Referral Hospital, Northwest Ethiopia: A case-control study. HIV AIDS (Auckl). 2017;9:153–159. https://doi.org/10.2147/HIV.S139516
- Rangarajan S, Colby DJ, Giang LT, et al. Factors associated with HIV viral load suppression on antiretroviral therapy in Vietnam. J Virus Erad. 2016;2(2):94–101. https://doi.org/10.1016/S2055-6640(20)30466-0
- Muri L, Gamell A, Ntamatungiro AJ, et al. Development of HIV drug resistance and therapeutic failure in children and adolescents in rural Tanzania: An emerging public health concern. Aids. 2017;31(1):61–70. https://doi.org/10.1097/ QAD.00000000001273
- Haghighat R, Toska E, Bungane N, Cluver L. The HIV care cascade for adolescents initiated on antiretroviral therapy in a health district of South Africa: A retrospective cohort study. BMC Infect Dis. 2021;21(1):1–9. https://doi. org/10.1186/s12879-020-05742-9
- Nabitaka VM, Nawaggi P, Campbell J, et al. High acceptability and viral suppression of patients on Dolutegravir-based first-line regimens in pilot sites in Uganda: A mixedmethods prospective cohort study. PLoS One. 2020;15(5):1–13. https://doi. org/10.1371/journal.pone.0232419
- Esber A, Dear N, Shah N, et al. Brief report: Virologic impact of the dolutegravir transition: Prospective results from the multinational african cohort study. J Acquir Immune Defic Syndr. 2022;91(3):285–289. https://doi.org/10.1097/ QAI.0000000000003065
- Barry O, Powell J, Renner L, et al. Effectiveness of first-line antiretroviral therapy and correlates of longitudinal changes in CD4 and viral load among HIV-infected children in Ghana. BMC Infect Dis. 2013;13(1). https://doi.org/10.1186/1471-2334-13-476

- 32. WHO. Updated recommendations on first-line and second-line antiretroviral regimens and post-exposure prophylaxis and recommendations on early infant diagnosis of HIV [homepage on the Internet]. WHO Guidelines; 2018 [cited 2024 Aug 25]. Available from: http://www.who.int/hiv/pub/guidelines/ ARV2018update/en/%0AWeb
- Devendra A, Kohler M, Letsika M, et al. HIV viral suppression in children and adolescents 2 years after transition to dolutegravir: A multicentre cohort study. AIDS. 2024;38(7):1013–1023. https://doi.org/10.1097/QAD.00000000003835
- Umeokonkwo CD, Onoka CA, Agu PA, Ossai EN, Balogun MS, Ogbonnaya LU. Retention in care and adherence to HIV and AIDS treatment in Anambra State Nigeria. BMC Infect Dis. 2019;19(1):1–11. https://doi.org/10.1186/s12879-019-4293-8
- Paterson DL, Swindells S, Mohr J, et al. Adherence to protease inhibitor therapy and outcomes in patients with HIV infection. Ann Intern Med. 2000;133(1):21–30. https://doi.org/10.7326/0003-4819-133-1-200007040-00004
- Sahay S, Srikanth Reddy K, Dhayarkar S. Optimizing adherence to antiretroviral therapy. Indian J Med Res. 2011;134(6):835–849. https://doi.org/10.4103/0971-5916.92629
- 37. Roscoe C, Hachey D. Retention in HIV care [homepage on the Internet]. Core Competencies for the National HIV Curriculum; 2022 [cited 2024 Aug 30]; p. 1–38. Available from: https://www.hiv.uw.edu/go/basic-primary-care/ retention-care/core-concept/all
- Mushy SE, Mtisi E, Mkawe S, et al. Barriers to viral load suppression among adolescents living with HIV on anti-retroviral therapy: A retrospective study in Tanga, Tanzania. AIDS Res Ther. 2024;21(1):1–8. https://doi.org/10.1186/s12981-024-00622-7
- Munyayi FK, Van Wyk B. Closing the HIV treatment gap for adolescents in Windhoek, Namibia: A retrospective analysis of predictors of viral nonsuppression. Int J Environ Res Public Health. 2022;19(22):14710. https://doi. org/10.3390/ijerph192214710
- 40. UNAIDS. Ending the AIDS epidemic for adolescents, with adolescents [homepage on the Internet]. 2016 [cited 2024 Sep 08]. Available from: https:// www.unaids.org/sites/default/files/media_asset/ending-AIDS-epidemicadolescents_en.pdf

- 41. Nyawira G, Samuel W, Maricianah O, Elizabeth AB, Kenneth N. Determinants of antiretroviral therapy adherence among older adolescents living with HIV in Kenya during the transition to adult care: An observational study. J AIDS HIV Res. 2020;12(2):24–33. https://doi.org/10.5897/JAHR2020.0513
- Van Wyk BE, Davids LAC. Challenges to HIV treatment adherence amongst adolescents in a low socio-economic setting in Cape Town. South Afr J HIV Med. 2019;20(1):1002. https://doi.org/10.4102/sajhivmed.v20i1.1002
- Van Wyk BE, Kriel E, Mukumbang FC. Two-year viral load suppression among adolescents receiving antiretroviral therapy in the Cape Metropole, South Africa, 2013 – 2015: A retrospective cohort analysis. S Afr Med J. 2020;110(12): 1213–1217. https://doi.org/10.7196/SAMJ.2020.v110112.14509
- 44. Orrell C, Hagins DP, Belonosova E, et al. Fixed-dose combination dolutegravir, abacavir, and lamivudine versus ritonavir-boosted atazanavir plus tenofovir disoproxil fumarate and emtricitabine in previously untreated women with HIV-1 infection (ARIA): Week 48 results from a randomised, open-label, non-inferiority, phase 3b study. Lancet HIV. 2017;4(12):e536–e546. https://doi.org/10.1093/ofid/ofw194.89
- 45. Van Lunzen J, Maggiolo F, Arribas JR, et al. Once daily dolutegravir (S/ GSK1349572) in combination therapy in antiretroviral-naive adults with HIV: Planned interim 48 week results from SPRING-1, a dose-ranging, randomised, phase 2b trial. Lancet Infect Dis. 2012;12(2):111–118. https://doi.org/10.1016/ S1473-3099(11)70290-0
- 46. Calmy A, Tovar Sanchez T, Kouanfack C, et al. Dolutegravir-based and low-dose efavirenz-based regimen for the initial treatment of HIV-1 infection (NAMSAL): Week 96 results from a two-group, multicentre, randomised, open label, phase 3 non-inferiority trial in Cameroon. Lancet HIV. 2020;7(10):e677–e687. https://doi. org/10.1016/S2352-3018(20)30238-1
- Sangeda RZ, Mosha F, Prosperi M, et al. Pharmacy refill adherence outperforms self-reported methods in predicting HIV therapy outcome in resource-limited settings. BMC Public Health. 2014;14(1):1035. https://doi.org/10.1186/1471-2458-14-1035
- Powell AE, Davies HTO, Thomson RG. Using routine comparative data to assess the quality of health care: Understanding and avoiding common pitfalls. Qual Saf Health Care. 2003;12(2):122–128. https://doi.org/10.1136/qhc.12.2.122