

# MINISTRY OF HEALTH OFFICE OF THE DIRECTOR GENERAL

Telephone:(020) 2717077 Fax: (020) 2713234

Email: <u>dghealth2019@gmail.com</u>

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Ref No: MOH/1/2/42

The Chief Executive Officer
Kenyatta National Hospital
NAIROBI

The Chief Executive Officer

Moi Teaching and Referral Hospital

ELDORET

The Chief Executive Officer
Mathari Hospital
NAIROBI

The Chief Executive Officer
Spinal Injury
NAIROBI

The Chief Executive Officer
Kenyatta University Teaching & Referral Hospital
OTHAYA

The Chief Executive Officer
Kenya HealthCare Federation (KHF)
admin@khf.co.ke

The Chief Executive Officer
Rural Private Hospitals Association of Kenya (RUPHA)
info@rupha.co.ke

AFYA HOUSE CATHEDRAL ROAD P. O. Box 30016 - 00100 NAIROBI

Date: 2<sup>nd</sup> January 2025

#### The Chief Executive Officer

Jaramogi Oginga Odinga Teaching and Referral Hospital **KISUMU** 

#### RE: OPTIMIZATION OF PLHIV ON ATV/R 300/100MG

Ministry of Health through National AIDS & STI Control Program (NASCOP) has over time provided guidance on treatment optimization for children, adolescents and adults living with HIV on Protease Inhibitors (PIs) in a phased manner putting into consideration research and commodity management needs. Listed below were the proposed phases for optimization of PLHIV on PIs:-

Phase 1: Optimization of Children and Adolescents Living with HIV (CALHIV) on first line PIs (Lopinavir/ritonavir, LPV/r) to DTG based regimens.

Phase 2: Optimization of Adults Living with HIV on First Line Protease Inhibitors (Lopinavir/ritonavir, LPV/r and Atazanavir/ritonavir, ATV/r) to TLD.

Phase 3: Optimization of Adults Living with HIV on 2nd Line ATV/r to DTG based regimens

Phase 4: Optimization of CALHIV on Second Line LPV/r to DTG based regimens

A committee of experts from NASCOP, KEMSA, County Governments, HIV Implementing Partners, technical experts from universities and community representatives reviewed the current HIV data from research and the commodity situation to inform the following guidance for adults and adolescents on ATV/r 300/100mg with immediate effect targeted for phase 3 i.e. Optimization of Adults Living with HIV on 2nd Line ATV/r to DTG based regimens.

## 1. PLHIV Currently on ATV/r 300/100mg who are virally suppressed (<1000 copies/mL)

a) INSII naive	Optimized ART Regimen	
Current Regimen	≤ 30kg	≥ 30kg
ABC/3TC+ATV/r	ABC/3TC+DTG	TDF/3TC/DTG(FDC)
AZT/3TC+ATV/r	AZT/3TC+DTG	TDF/3TC/DTG(FDC)
TDF/3TC+ATV/r		TDF/3TC/DTG(FDC)

### b) INSTI experienced

Maintain on ATV/r 300/100mg but consider transition to a superior in class PI (DRV/r 400/50mg) once available.

# 2. PLHIV on ATV/r 300/100mg who are not virally suppressed (>1000 copies/mL)

### a) Both INSTI Naïve and INSTI experienced

Maintain on ATV/r 300/100mg, address any adherence issues by instituting the interventions for Suspected treatment failure as per ART guidelines 2022, once virally suppressed transition as above. For those with confirmed treatment failure, collect a DRT sample and switch based on the DRT results. However, given the stock out of DRT reagents, collect the DRT sample and then switch as guided below until when the results are available to guide the appropriate regimen.

	Optimized ART Regimen	
<b>Current Regimen</b>	≤ 30kg	≥ 30kg
ABC/3TC+ATV/r	Switch to AZT/3TC+DTG	Switch to TDF/3TC/DTG
AZT/3TC+ATV/r	Switch to ABC/3TC+DTG	Switch to TDF/3TC/DTG
TDF/3TC+ATV/r	•	Switch to TDF/3TC/DTG

### 3. PLHIV with confirmed treatment failure on TLD as first line.

Switch to a PI based second line regimen with ATV/r 300/100mg as a holding second line and transition to DRV/r 400/50mg as preferred second line once available.

#### NOTE

- PLHIV on dual regimen containing ATV/r 300/100mg will be retained on their regimen until a superior in class PI (DRV/r 400/50mg) is available
- PLHIV on third line regimens consisting of ATV/r 300/100mg will be transitioned on a case to case basis in consultation with the Regional TWGs.
- To avert any wastages, ART pharmacy managers should ensure the existing stocks of ATV/r are fully utilized or planned for utilization (for clients who may still require ATV/r) prior to transition of eligible clients.

• The optimization should be carried out at the scheduled appointments for clients hence there should be no active recall of clients to health facilities.

For any further clarification kindly reach out to Dr. Rose Wafula at <a href="mailto:head@nascop.or.ke">head@nascop.or.ke</a> and copy <a href="mailto:carentreatmentmanager@nascop.or.ke">carentreatmentmanager@nascop.or.ke</a>.

Dr. Patrick Amoth, EBS

**DIRECTOR GENERAL FOR HEALTH**