# Early effect of transitioning children to dolutegravir-based regimens on viral load suppression in Malawi

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### BACKGROUND

- Viral suppression (VS) in children has remained suboptimal compared to adults.
- In 2021, EGPAF and the Ministry of Health in Malawi rolled out a pediatric formulation of dolutegravir (10mg film-coated tablet) (pDTG) in children <20 kgs.
- We evaluated the impact of transitioning children <20 kgs to pDTG on VS in Malawi.

### **METHODS**

- We analyzed routine retrospective program data from electronic medical record systems (EMRS) pooled across 169 healthcare facilities in Malawi supported by the Elizabeth Glaser Pediatric AIDS Foundation (EGPAF) in eight districts.
- Children under 15 years and weighing less than 20 kgs, receiving antiretroviral therapy (ART) between July 2021 and June 2022, were included.
- We described the distribution of demographic and clinical characteristics, ART regimen, ART adherence (good adherence defined as missing no more than two antiretroviral doses in a month at the last follow-up visit), and VS (<1000 copies/mL).
- We used a logistic regression to identify factors associated with post-transition VS, adjusting for demographic characteristics, initial ART regimen, guardian type, adherence, and pre-transition viral load.

## Main Findings

38.6% (n=860) of children had routine viral load testing results six months following the transition to pDTG-based ART; of those with a viral load test, 81.1% (n=700) achieved VS.

#### RESULTS

- 2,468 children living with HIV (CLHIV) were included, 55.3% (n=1,364) of whom were age <60 months.
- 90.4% (n=2,230) had initiated non-DTG-based ART before pDTG was available.

Table 1: Factors associated with viral suppression among CLHIV <20 kgs, receiving ART in EGPAFsupported facilities, who transitioned to pDTG-based ART (Logistic regression)

Characteristic	VL <1000 copies/m	L Unadjusted ORs	Adjusted ORs
	n (%)	(95% CI)	(95% CI)
Sex			
Female	394 (81.2%)	Ref	_
Male	306 (81.0%)	0.98 (0.70-1.38)	1.03 (0.62-1.71)
Age range (months)			
<12	5 (62.5%)	Ref	_
12-23	41 (75.9%)	1.89 (0.40-9.02)	_
24-59	294 (78.2%)	2.15 (0.50-9.19)	_
60+	360 (84.7%)	3.32 (0.78-14.25)	_
<b>Facility location</b>			
Urban	253 (83.2%)	Ref	_
Rural	447 (80.0%)	0.80 (0.56-1.16)	_
EGPAF clinical staff			
available on site			
Yes	500 (82.1%)	Ref	_
Νο	200 (78.7%)	0.81 (0.56-1.16)	_
Initial ART Regimen			
PI-Based*	549 (81.7%)	Ref	1
NNRTI-Based**	95 (76.6%)	0.73 (0.46-1.16)	-
Other	3 (60.0%)	0.34 (0.05-2.03)	-
Adherence to ARVs			
<b>Poor Adherence</b>	281 (75.3%)	Ref	1
<b>Good Adherence</b>	419 (85.5%)	1.93 (1.37-2.73)	2.79 (1.65-4.71)
<b>Pre-transition Viral</b>			
load			

- 62.7% (n=1,398) of the patients who had been on other ART regimens, had a viral load (VL) test result before the transition to pDTG; of those tested, 62.1% (n=868) achieved VS.
- 99.9% of CLHIV (n=2,227) transitioned to pDTG-based regimens (without change in nucleoside backbone). 52.9% (n=1,179) of them had good ART adherence at 6 months posttransition.
- 38.6% (n=860) had routine VL testing results six months following the transition to pDTGbased ART; of those tested,

81.1% (n=700) achieved VS.

Good ART adherence and being virally suppressed before transition was associated with post-transition VS

≥1000 copies/mL 181 (71.2%) Ref <1000 copies/mL 5.53 (3.23-9.48) 358 (93.0%) 5.32 (3.30-8.57)

\*Protease inhibitor-based, \*\*Non-Nucleoside reverse transcriptase inhibitor-based **CONCLUSIONS** 

- VS was achieved in most children tested within the first six months after pDTG transition.
- However, adherence was suboptimal in this group, and VL testing at six months posttransition was limited.
- Interventions to improve VL testing and enhance good adherence are needed in children to continue progressing towards the 95-95-95 UNAIDS goals.

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