

# Clinical outcomes and renal safety in HIV/AIDS patients on tenofovir-containing regimens in Lesotho

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## BACKGROUND

- Tenofovir (TDF)-containing highly active antiretroviral (HAART) regimens are the most preferred in the treatment of HIV/AIDS in Lesotho
- Tenofovir (TDF) inclusion in HAART regimens has clinically shown to have an excellent efficacy and safety outcomes when compared with other HAART regimens containing other first-line antiretroviral drugs, such as abacavir (ABC), stavudine (d4T) and zidovudine (AZT).<sup>45</sup>
- Although TDF use has been associated with acceptable safety, several studies have reported a rare manifestation of renal disease in HAART regimens that include TDF.<sup>123678</sup>

## PROJECT AIM

To evaluate the clinical outcomes (weight and CD4 count) and renal safety (eGFR) in HIV/AIDS patients taking TDF-containing HAART regimens in Lesotho

## METHODS

Descriptive, observational, longitudinal retrospective design was followed on 255 adult patients treated on TDF-containing HAART regimens at primary health care facility called Paballong HIV/AIDS care centre located in Berea district in Lesotho; from October 2015 to march 2016.

## RESULTS

Body weight according to treatment duration, sex and age at antiretroviral therapy initiation: Estimates of fixed effects							
Parameter	Estimate	Std. Error	Df	t	p value	95% Confidence Interval	
						Lower Bound	Upper Bound
Intercept	56.57	0.80	5414.89	70.75	0.00	55.10	58.13
Treatment duration	-0.003	0.002	188.66	-1.26	0.21	-0.01	0.002
[Sex=female]	2.64	0.39	5413.30	6.83	0.00	1.89	3.40
[Sex=male]	0	0					
Age at ART initiation	0.10	0.017	5414.39	5.52	0.00	0.06	0.13

CD4 cell count according to treatment duration, sex and age at antiretroviral therapy initiation: Estimates of fixed effects							
Parameter	Estimate	Std. Error	df	T	p value	95% Confidence Interval	
						Lower Bound	Upper Bound
Intercept	179.67	21.58	1585.19	8.33	0.00	137.33	222.10
Treatment duration	0.20	0.03	103.01	7.62	0.00	0.15	0.27
[Sex=female]	69.13	10.35	1588.70	6.68	0.00	48.83	89.43
[Sex=male]	0	0					
Age at antiretroviral therapy initiation	1.32	0.47	1591.99	2.78	0.00	0.39	2.25

Estimated glomerular filtration rate according to treatment duration, sex, age at antiretroviral therapy initiation and body weight: Estimates of fixed effects							
Parameter	Estimate	Std. Error	df	t	p value	95% Confidence Interval	
						Lower Bound	Upper Bound
Intercept	143.84	7.17	923.10	20.07	0.00	129.77	157.91
Treatment duration	0.000	0.003	153.24	0.23	0.82	-0.01	0.007
[Sex=female]	-13.05	2.36	900.44	-5.54	0.00	-17.67	-8.43
[Sex=male]	0	0					
Age at antiretroviral therapy initiation	-0.78	0.10	887.00	-7.50	0.00	-0.99	-0.58
Body weight	-0.02	0.09	903.84	-0.25	0.80	-0.20	0.16

## CONCLUSIONS

- Clinical outcomes manifesting by weight gain and CD4 cell count elevation improve upon initiation of antiretroviral therapy at any age.
- Females are far more at advantage to experience better clinical outcomes than males over the treatment duration.
- The renal function is progressively deteriorated following initiation of antiretroviral therapy at any age.
- Females experience more renal compromise than males over the treatment duration.

### Reference list

- Chua, A.C., Llorin, R. M., Lai, K., Cavaller, P. & Law, H.L. 2012. Renal safety of tenofovir containing antiretroviral regimen in a Singapore cohort. *AIDS research and therapy*, 9:1-5.
- Cooper, R.D., Wiebe, N., Smith, N., Keisor, P., Naicker, S. & Tonelli, M. 2010. Systematic review and meta-analysis: renal safety of tenofovir disoproxil fumarate in HIV-Infected patients. *Clinical infectious diseases*, 51:496-505.
- Crum-Cianflone, N., Ganesan, A., Teneza-Mora, N., Riddle, M., Medina, S., Barahona, I. & Brodine, S. 2010. Prevalence and factors associated with renal dysfunction among HIV-infected patients. *AIDS patient care and STDs*, 24(6):353-360.
- Gallant, J.E., DeJesus, E., Arribas, J.R., Pozniak, A.L., Gazzard, B., Campo, R.E., Lu, B., McColl, D., Chuck, S., Enejosa, J., Toole, J.J. & Cheng, A.K. 2006. Tenofovir DF, emtricitabine, and efavirenz vs. zidovudine, lamivudine, and efavirenz for HIV. *The new England journal of medicine*, 324(3):251-260.
- Gallant, J.E., Staszewski, S., Pozniak, A.L., DeJesus, E., Suleiman, J.M.A.H., Miller, M.D., Coakley, D.F., Lu, B., Toole, J.J. & Cheng, A.K. 2004. Efficacy and safety of tenofovir DF vs stavudine in combination therapy in antiretroviral-naïve patients: a 3-year randomized trial. *Journal of the American Medical Association*, 292(2):191-201.
- Johnson, D.C., Chasela, C., Maliwichi, M., Mwafongo, A., Akinkoutu, A., Moses, A., Jamieson, D.J., Kourtis, A.P., King, C.C., Horst, C. & Hosseinipour, M.C. 2012. Tenofovir use and renal insufficiency among pregnant and general adult population of HIV-infected, ART-naïve individuals in Lilongwe, Malawi. *Public library of science one*, 7(7):1-7.
- Kalysesubula, R. & Perazella, M.A. 2011. Nephrotoxicity of HAART. *Hindawi*: 1-11.
- Sadre, A., Munshi, N., Dhande, S. & Dravid, A. 2012. Tenofovir-induced acute kidney injury in HIV-infected patients in western India: a resource limited setting perspective. *Journal of the International AIDS Society*, 15:1.