Favorable metabolic outcomes 48 weeks after switch to DTG/3TC

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BACKGROUND

New agents (more potent) Therapy similaritication Procommendation of dual ART

Recommendation of 2DR
paradigm shift in HIV treatment

 2nd generation integrase inhibitors and tenofovir alafenamide (TAF) have been associated with weight gain

→ impact on metabolic health and cardiovascular risk?



METHODS

Randomized, open-label controlled trial (2:1)

Switch to DTG/3TC Switch or stay on BIC/FTC/TAF

- Longitudinal follow-up: baseline, week 24, week 48
- Outcomes (2ary): weight, BMI, waist, lipids, insulin resistance, DXA scan, fibroscan
- Linear mixed models with covariance patterns

• Intention to treat – exposed analysis



Contact: Sophie Degroote, PhD Algemene Inwendige Ziekten, UZ Gent Corneel Heymanslaan 10, 9000 Gent (Belgium) T +32 (0)9332 01 55 / E sophie.degroote@uzgent.be We thank our supporting partner, ViiV Healthcare. Switch to DTG/3TC may have a favorable impact on metabolic outcomes at week 48 as compared to BIC/FTC/TAF.

RESULTS

 Weight, waist and BMI were different at baseline between both treatment groups.

Baseline characteristics	DTG/3TC (n = 87)	BIC/FTC/TAF (n = 43)	p-value
Male sex (%)	90,8	90,7	1.000
Age (mean ± SD)	47,3 ± 11,9	45,0 ± 11,6	0.292
Non-European ethnicity (%)	19,5	25,6	0.628
Sexual orientation (%)			0.526
Gay/lesbian or bisexual/pansexual	70,1	67,4	
Heterosexual	27,6	30,2	
ART regimen at baseline (%)			0.072
DTG/ABC/3TC	31	51	
BIC/FTC/TAF	68	49	
Years on ART (median (IQR))	8 (5-11)	6 (4-9)	0.133
Years on 2nd generation INSTI (median (IQR))	3 (2-5)	4 (3-5)	0.476
CD4 nadir (cells/µl; median (IQR))	206 (193-476)	269 (212-380)	0.510
Weight (kg; mean ± SD)	81 ± 12	75 ± 13	0.013
Waist (cm; mean ± SD)	95 ± 12	89 ± 11	0.006
BMI (kg/m²; median (IQR))	26 (23-28)	25 (22-26)	0.024

 Linear mixed models (adjusted for baseline BMI) revealed significantly different estimated mean differences from baseline to week 48 between both groups with regard to ALT, HDL cholesterol, lean trunk mass, trunk fat mass and fat percentage.

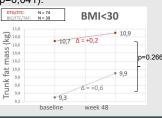
There were no significant differences with regard to the other outcomes (lean body mass,

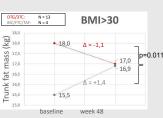
glucose, insulin, noma-in, iver horo				
	DTG/3TC	BIC/FTC/TAF		
ALT (U/L)	- 0.73	+ 4.55		
HDL (mg/L)	- 0.043	- 2.84		
Lean trunk mass (gram)	+ 112	- 474		
Trunk fat mass (gram)	+ 41	+ 719		
Fat percentage	- 0.04	+ 1.32		

total and LDL cholesterol, triglycerides,

	DTG/3TC	BIC/FTC/TAF	p-value
Weight (kg)	+0,29	+0,30	0.987
Waist (cm)	-0,07	+ 1,10	0.155
BMI (kg/m ²)*	+0,07	+0,04	0.919
Cholesterol (mg/dl)	-2,49	-8,90	0.316
LDL cholesterol (mg/dl)	-1,82	-6,21	0.435
TC/HDL	0	+0,029	0,848
Triglycerides (mg/dl)	-3,82	-20,96	0.206
HOMA-IR	-0,16	-0,43	0.359
FibroCAP (dB/m)	-0,39	-11,61	0.304

 Greater treatment-mediated differences in trunk fat were observed in people with BMI > 30 (p=0.041).







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DISCUSSION

- Viral load suppression maintained throughout study, without differences between groups
- Importance of body composition measures, in addition to weight

Limitations

- Small study sample / Singe center study
- Baseline differences
- Woman & sub-Saharan Africa people underrepresented
- Lifestyle parameters not measures

Future directions

- Further longitudinal data (follow-up until week 144)
- · Virological and inflammatory analyses to be done