

0980 TFV-DP in DBS for pregnant/postpartum adolescent and young women on PrEP in Africa

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BACKGROUND

- The risk of HIV acquisition more than doubles in pregnant and postpartum women in sub-Saharan Africa, emphasizing the need for programmatic delivery of HIV prevention services including PrEP.
- Intracellular tenofovir-diphosphate (TFV-DP) in red blood cells measured with dried blood spots (DBS) is an established biomarker of cumulative PrEP adherence given its 17 day half-life, analogous to HbA1C for diabetes.
- TFV-DP in DBS informs adherence-efficacy relationships for PrEP. This has been defined for men who have sex with men, but not for pregnant or non-pregnant women.
- Pregnancy causes physiological changes expected to lower TFV-DP in DBS including increased TFV renal clearance and hemodilution.
- IMPAACT 2009 is a two-component observational trial in pregnant and postpartum adolescent girls and young women in sub-Saharan Africa. This communication presents results from the first (PK) component.

OBJECTIVES

The goal of the IMPAACT 2009 PK component was to establish adherence benchmarks for TFV-DP in DBS for pregnant and postpartum adolescents and young women who took PrEP daily under direct observation, and to compare these benchmarks in the pregnant and postpartum groups.

METHODS

- HIV-negative adolescent girls and young women (16-24 years) were recruited in Malawi, South Africa, Uganda, and Zimbabwe
- Pregnant: 14-24 wks gestation at enrollment
- Postpartum: 6-12 wks postpartum at enrollment
- Daily FTC/TDF was administered for 12 weeks under direct observation (in person or by live video streaming).
- Five 50uL DBS were collected weekly and stored at -80°C.
- TFV-DP was assayed in one 50uL spot by validated LC-MS/MS at the University of Cape Town and reported as fmol/3mm punch for consistency with previous studies.
- Summary statistics were used for observed data at week 12.
- A one-compartment IV infusion non-linear mixed effects model was fit to concentration time data for individual predictions (modeled data).
- Observed TFV-DP at week 12 was compared between the pregnant and postpartum groups with the Wilcoxon test.

RESULTS

- 20 pregnant and 20 postpartum women enrolled between March-May 2019
- 3348 of 3360 (>99%) total doses were directly observed
- All participants met criteria for inclusion in the analysis

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Adherence benchmarks using TFV-DP in DBS were established for pregnant/postpartum African adolescents and young women

TFV-DP in DBS was 31%-37% lower in pregnancy compared with postpartum, in line with expectations. Strict adherence to PrEP is recommended during pregnancy.

Table 1. Baseline Demographics

Characteristic	Pregnant (N=20) median (IQR)	Postpartum (N=20) median (IQR)
Age (yrs)	20 (19.5, 22.5)	20 (19, 22)
Weight (kg)	59 (56, 65)	55 (51, 62)
HCT (%)	34.9 (33, 37.1)	40.8 (39.1, 41.7)
CLcr (CG-ideal wt; ml/min)	151 (130, 169)	109 (102, 123)
Gestational Age (wks)	18 (15, 20)	NA
Postpartum (wks)	NA	7 (7, 9)

Table 2. TFV-DP at Week 12 Visit

Characteristic	Pregnant (N=20) median (IQR)	Postpartum (N=20)* median (IQR)	Difference (p Wilcoxon)
Observed TFV-DP (fmol/punch)	965 (691, 1166)	1406 (1053, 1859)	31% (p=0.0064)
Modeled TFV-DP (fmol/punch)	890 (704, 1143)	1418 (1179, 2139)	37% (p<0.0001)
Modeled T-1/2 (days)	14 (10.6, 17.6)	16.5 (13.7, 21.2)	ND

One week 12 value was missing, and their concentration at week 11 was used. Median IQR was 1434 (966, 1869) if only week 12 was used.

- Figures 1 and 2 show TFV-DP concentration time profiles for observed (Figure 1) and modeled (Figure 2) data.
- Figure 3 shows estimated thresholds. These were based on 25th percentiles of observed categories were determined assuming dose-proportionality.

data that were rounded down to establish the highest adherence group. Lower adherence







- Adherence established

The authors would like to thank the study participants, study staff, the University of Cape Town laboratory, Mustafa Ibrahim and Jenna Yager for modeling contributions. Overall support for IMPAACT was provided by NIAID with co-funding from NICHD and NIMH, under Award Numbers UM1AI068632, UM1AI068616 and UM1AI106716, and NICHD contract number HHSN2752018000011. The content is solely the responsibility of the authors and does not necessarily represent the official views of the NIH. Study drug was provided by Gilead Sciences.

Figure 2. Modeled TFV-DP

CONCLUSIONS

benchmarks DBS for TFV-DP in for pregnant and postpartum adolescents and young women (Figure 3, right).

• TFV-DP in DBS was 31%-37% lower in pregnant compared with postpartum women, in parallel with increased renal function and lower HCT. This is consistent with prior studies.

• Strict PrEP adherence is recommended during pregnancy. Future studies can use these adherence benchmarks to support PrEP adherence and determine adherence-efficacy relationships.

were Figure 3. Proposed thresholds DBS TFV-DP fmol/punch

Interpretation	Pregnant	Post-partum
~7 doses/wk	≥650	≥950
2–6 doses/wk	200-649	250-949
<2 doses/wk	<200	<250

ACKNOWLEDGEMENTS