

MTN-017 Update

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MTN-017

- A Phase 2 Randomized Sequence Open Label Expanded Safety and Acceptability Study of Oral Emtricitabine/Tenofovir Disoproxil Fumarate Tablet and Rectally-Applied Tenofovir Reduced-Glycerin 1% Gel

Study Products

□ TDF/FTC



□ 1% tenofovir RG Gel



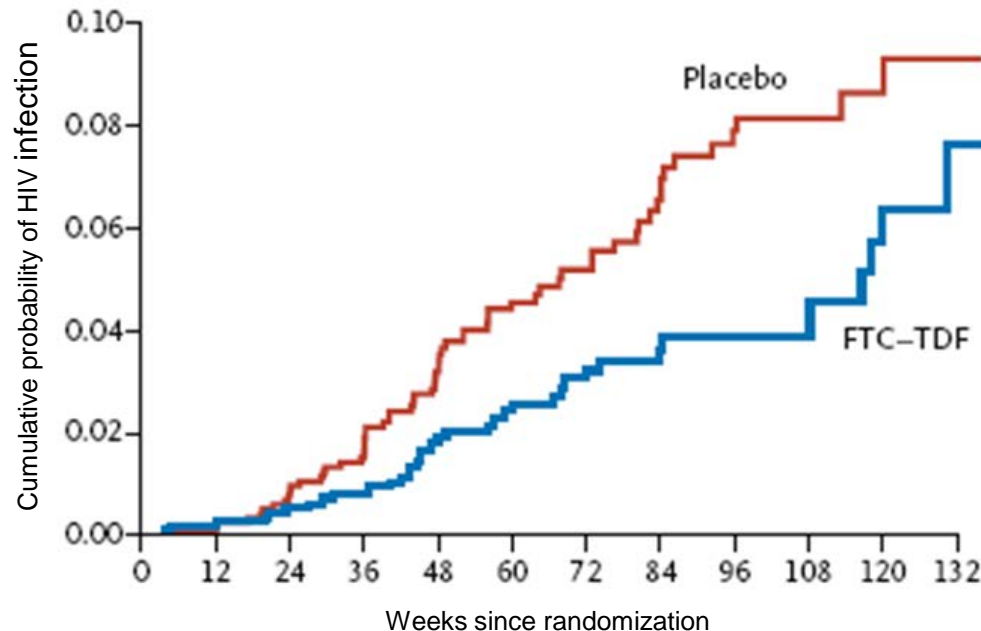


iPrEx Study

- Phase III multicenter trial of 2499 HIV negative MSM or transgender women randomized to receive placebo vs. FTC/TDF
- Median follow up 1.2 years
- Primary Endpoints:
 - AEs
 - HIV seroconversion

iPrEx: HIV Seroconversion

- 44% effectiveness
(95% confidence interval, 15 to 63; $P=0.005$)
- ~50% adherence to product

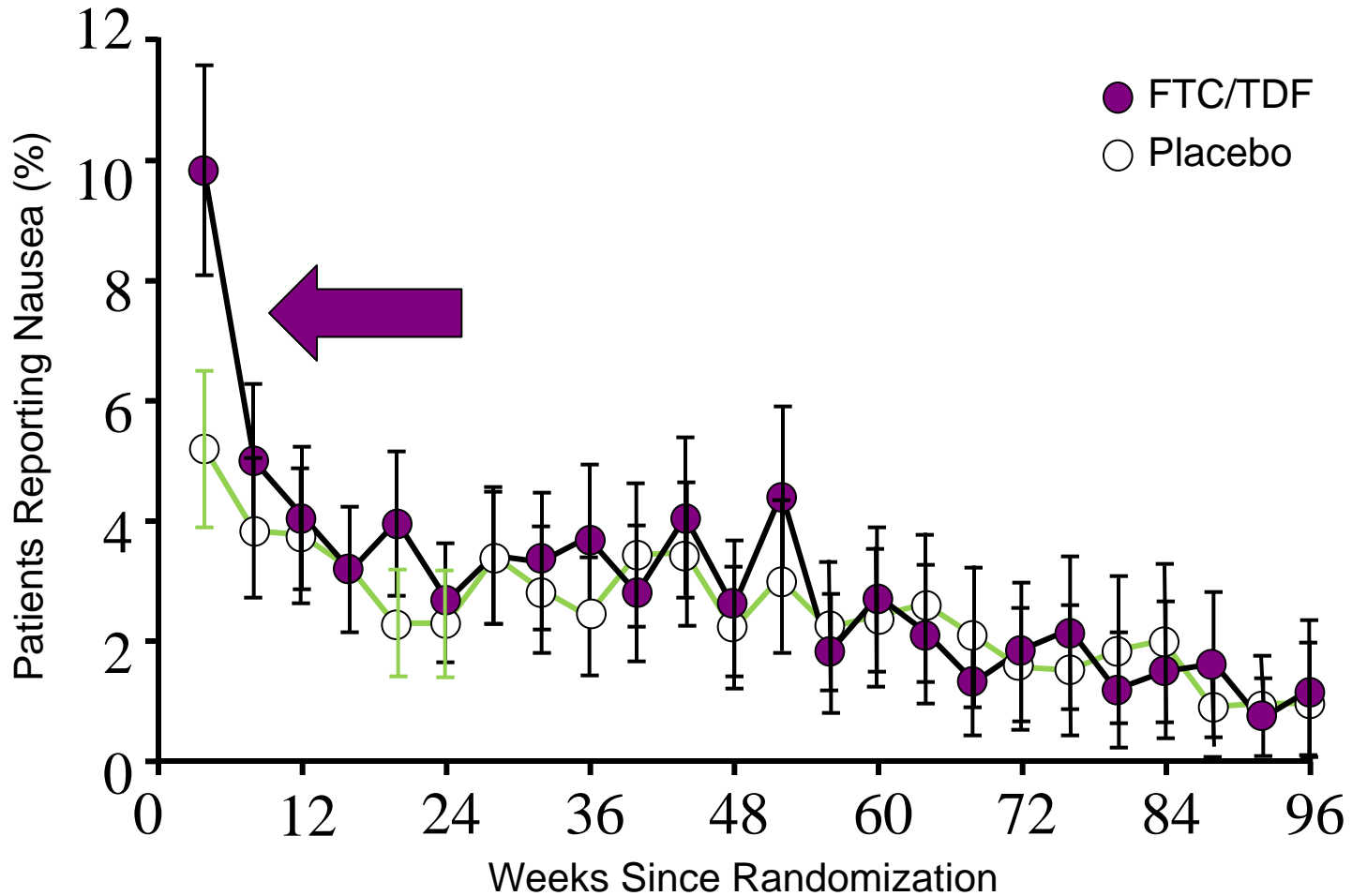


iPrEx: Adverse Events

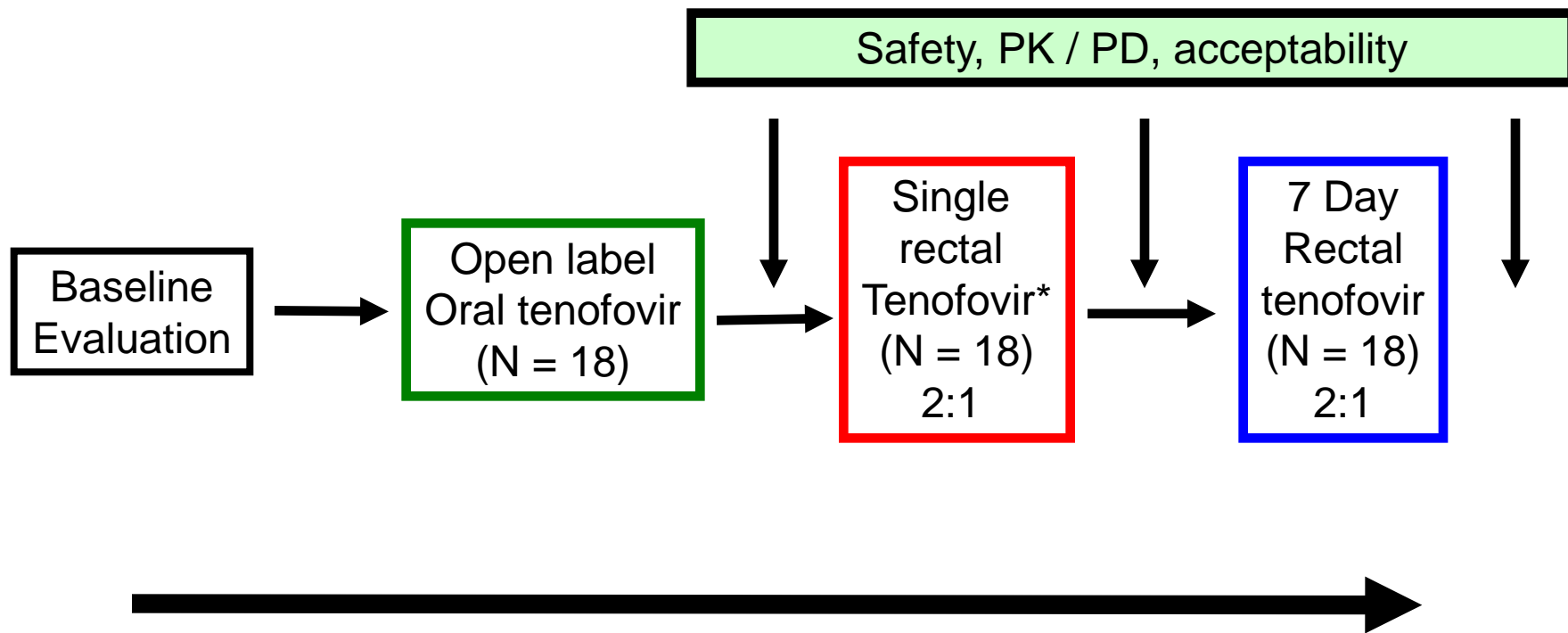
- No significant differences between active and placebo arms for:
 - Any grade 3/4 event, death, SAE, elevated creatinine, creatinine elevation confirmed on next visit

Adverse Event	FTC/TDF (n = 1251)		Placebo (n = 1248)		P Value
	%	Events	%	Events	
Nausea	2	22	<1	10	0.04
Weight decrease	2	34	1	19	0.04

iPrEx: Nausea



RMP-02/MTN-006 Study Design

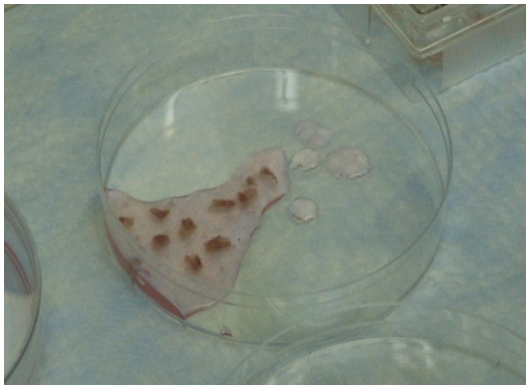


*1% tenofovir vaginal formulation

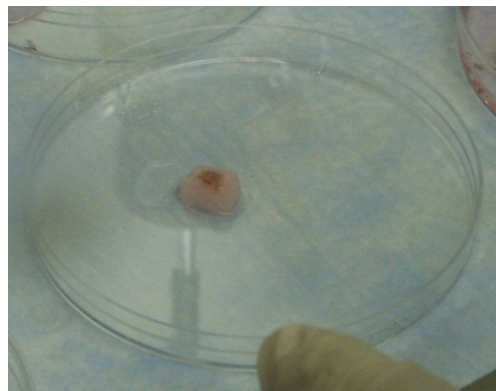
RMP-02/MTN-006 Adverse Events

GI Adverse Events in the Tenofovir Arm	RMP-02/MTN-006 (N = 12) Vaginal Formulation	
	N	%
Abdominal pain	6	50%
Rectal urgency	5	42%
Bloating	5	42%
Nausea	4	33%
Diarrhea	7	58%
Flatulence	3	25%
Proctalgia	0	0%
Other	5	42%
Total	35	-

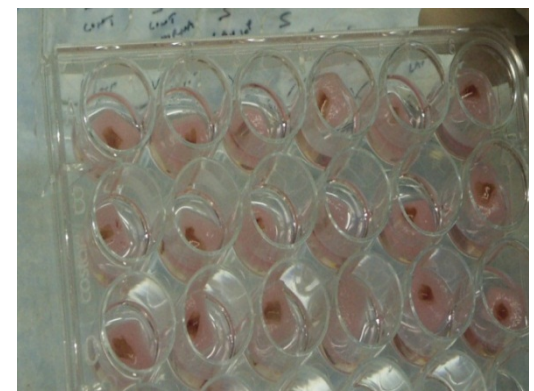
Colorectal Explants



Collect rectal biopsies
From ppts previously
exposed to Tenofovir gel



Place biopsy on raft

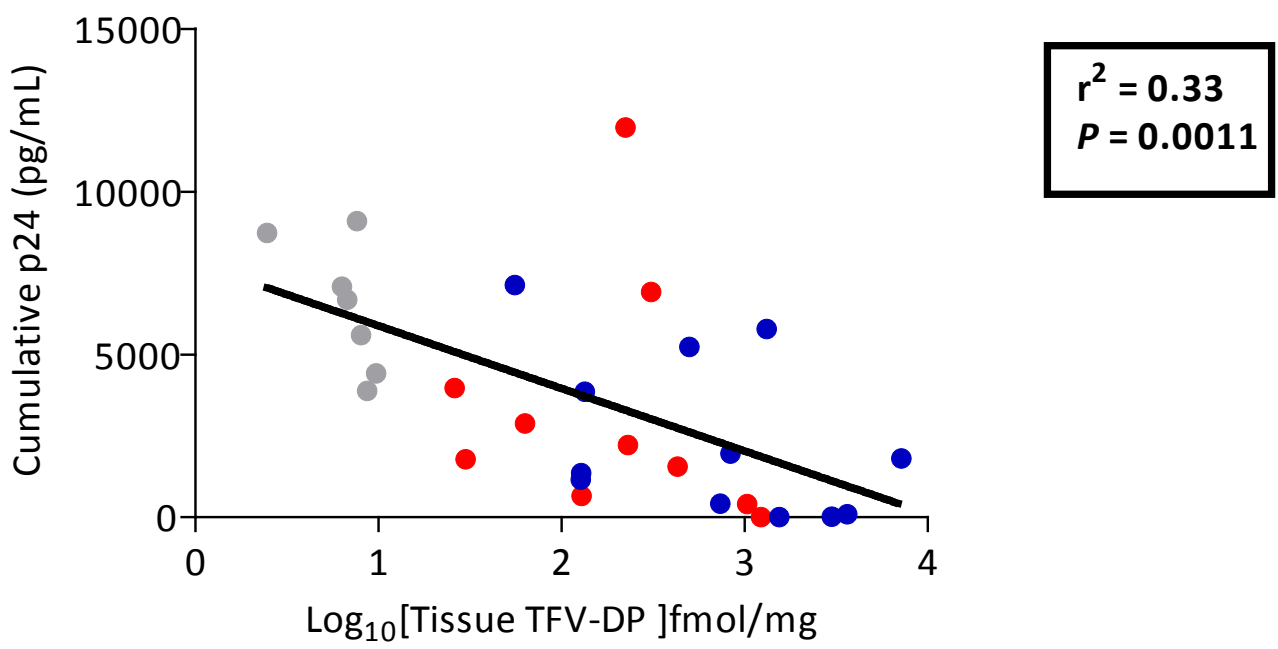


Expose to HIV and
measure sequential
p24 levels



PK/PD Correlation in RMP-02/MTN 006

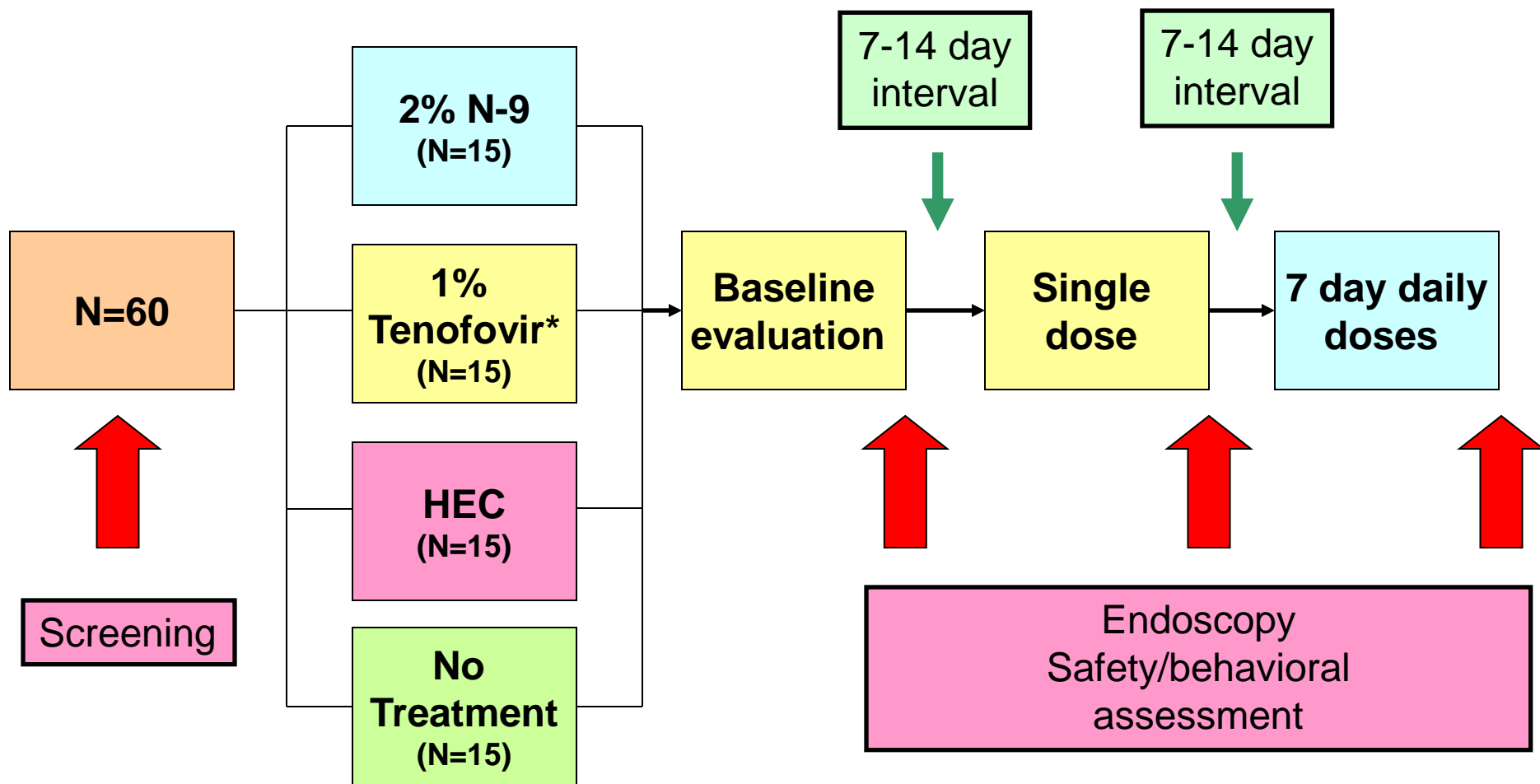
● Oral Dose ● Single Rectal Dose ● Multiple Rectal Dose



Acceptability

Product (N)	Like very much (%)
Tenofovir 1% gel (12)	25%
HEC Placebo (6)	50%

MTN-007 Study Design



*1% tenofovir reduced glycerin formulation

MTN-007 Adverse Events

GI Adverse Events (Tenofovir Arm)	MTN-007 (N = 16) RG Formulation	
	N	%
Abdominal pain	3	16%
Rectal urgency	0	0%
Bloating	0	0%
Nausea	0	0%
Diarrhea	1	6%
Flatulence	6	38%
Proctalgia	1	6%
Other	4	25%
Total	15	-

Gastrointestinal Adverse Events

GI Adverse Events in the Tenofovir Arm	MTN-007 (N = 16) RG Formulation		RMP-02/MTN-006 (N = 12) Original Formulation	
	N	%	N	%
Abdominal pain	3	16%	6	50%
Rectal urgency	0	0%	5	42%
Bloating	0	0%	5	42%
Nausea	0	0%	4	33%
Diarrhea	1	6%	7	58%
Flatulence	6	38%	3	25%
Proctalgia	1	6%	0	0%
Other	4	25%	5	42%

Acceptability

Product (N)	Intention to Use (%)
RG Tenofovir (15)	87%
HEC Placebo (15)	93%
N-9 (16)	63%



MTN 017

□ Study Population

- Approximately 186 pts
- HIV-uninfected
- MSM or transgender females
- Reported practicing receptive anal intercourse
- Age 18 years or older

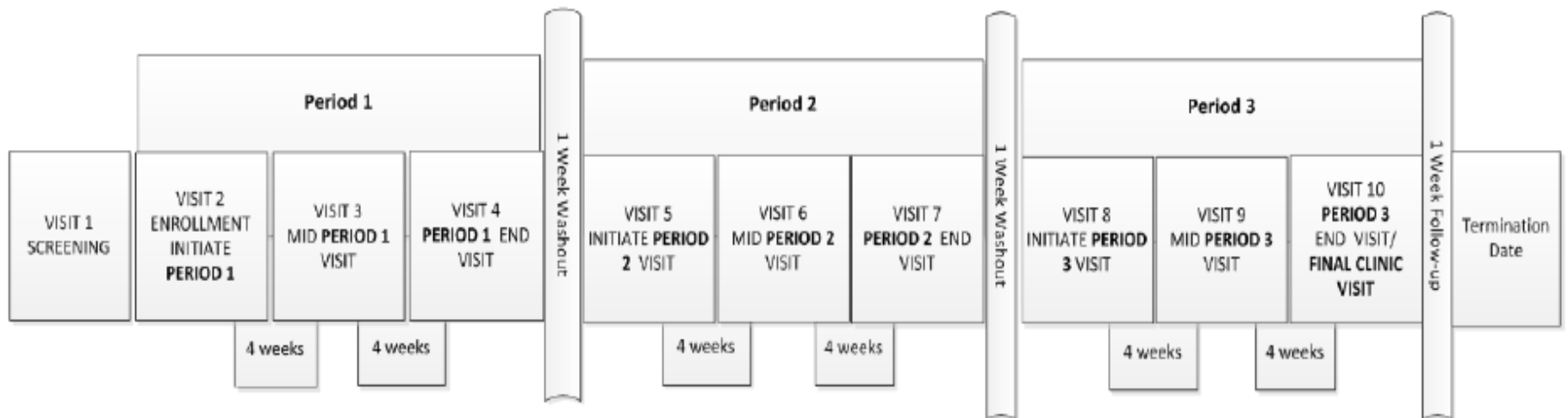
□ Study Duration

- Follow-up: 27 weeks per participant
- Accrual: Projected 6-9 calendar months at each site

MTN-017

- Study regimens include:
 - Rectal tenofovir gel used daily
 - Rectal tenofovir gel used before and after sex (BAT 24)
 - TDF/FTC tablets taken daily
- Each participant will follow all of the study regimens for eight weeks, with a weeklong break between regimens when no product will be used
 - The order in which participants follow study regimens will be based on random assignment
- All participants will receive standard HIV prevention package

MTN-017 Study Design



MTN-017 Study Design

Product Sequence	N	Period 1 (8 weeks)	Product Break (1 week)	Period 2 (8 weeks)	Product Break (1 week)	Period 3 (8 weeks)
1	31	Daily TDF/FTC		Daily rectal gel		Rectal gel before and after sex
2	31	Rectal gel before and after sex		Daily TDF/FTC		Daily rectal gel
3	31	Daily rectal gel		Rectal gel before and after sex		Daily TDF/FTC
4	31	Daily rectal gel		Daily TDF/FTC		Rectal gel before and after sex
5	31	Daily TDF/FTC		Rectal gel before and after sex		Daily rectal gel
6	31	Rectal gel before and after sex		Daily Rectal gel		Daily TDF/FTC

Study Sites





Primary Objectives/endpoints

□ Safety

- Compare the safety profiles of rectal tenofovir gel used daily and before and after sex, and TDF/FTC tablets
 - Grade 2 or higher adverse events

□ Acceptability

- To evaluate and compare acceptability of rectal tenofovir gel used daily and before and after sex, and TDF/FTC tablets
 - Participant self-report of ease of use, liking the product, and likelihood of product use if shown to be effective

Secondary Objective/Endpoints #1

□ Pharmacokinetics

- To compare systemic and local pharmacokinetics
 - Tenofovir concentrations
 - blood plasma, rectal tissue* and rectal fluid
 - Tenofovir-diphosphate concentrations
 - peripheral blood mononuclear cell (PBMC) and rectal tissue*
 - Emtricitabine concentrations
 - blood plasma, rectal tissue* and rectal fluid
 - Emtricitabine-triphosphate concentrations
 - PBMC and rectal tissue*

* Rectal tissue will be collected on a subset of participants taking part in the Rectal Biopsy/Fluid Subset



Secondary Objective/Endpoint #2

- Adherence
 - Compare the safety profiles of rectal tenofovir gel used daily and before and after sex, and TDF/FTC tablets
 - Percentage of prescribed doses taken orally or administered rectally in an 8-week period



Exploratory Objectives

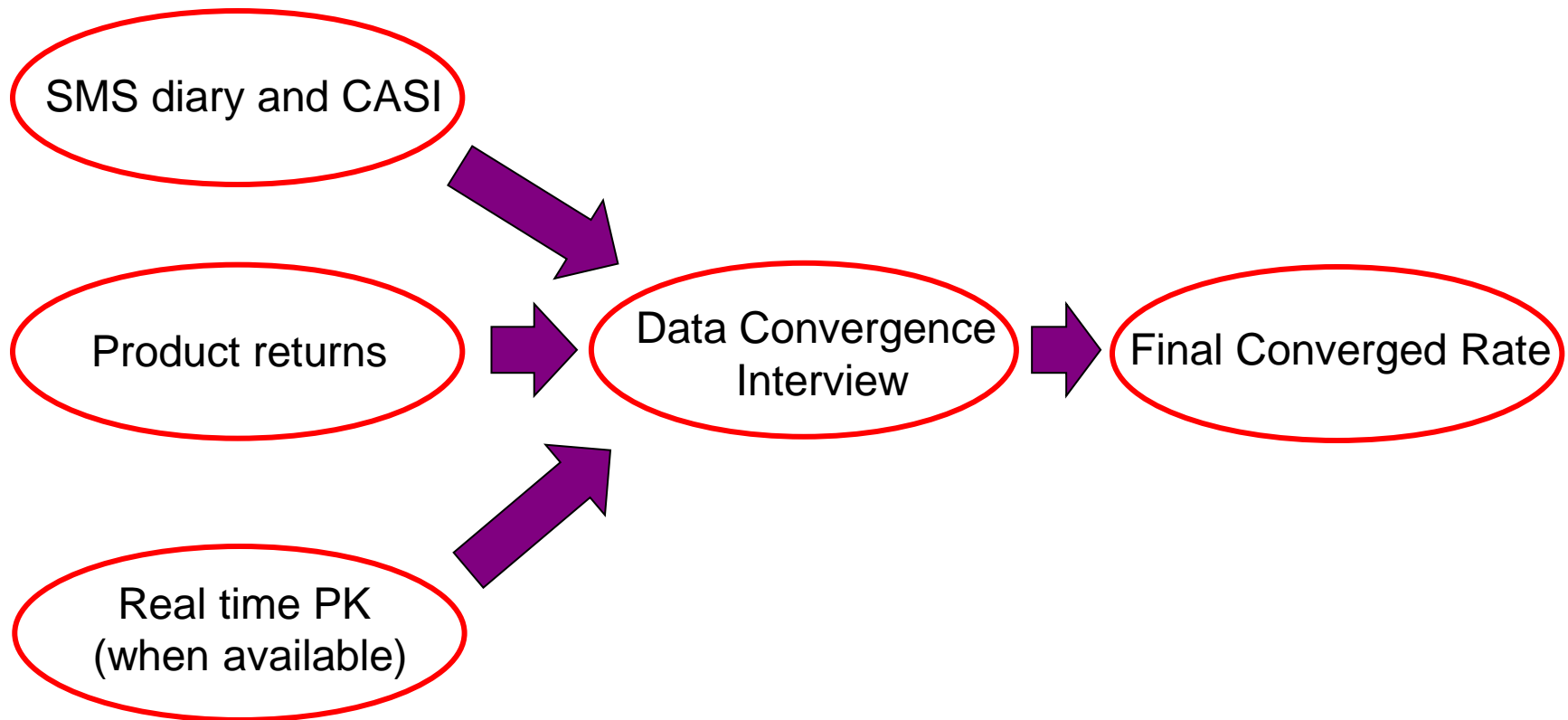
- Characterize pharmacodynamic responses
- Characterize changes in mucosal immunity
- Assess correlation between PK and adherence measures
- Identify factors associated with product adherence and whether they differ by product used
- Examine whether sexual activity or condom use varies by product used
- Determine the level of sharing of study products with non-participants
- Determine the prevalence of behavioral practices associated with anal intercourse that may affect microbicide use

Adherence in African PrEP Trials

Name	Population	Estimated Adherence		
		Self report	CPC	Drug level (in subset)
TDF ₂	557 ♀ & 662 ♂	94%	84%	80%
Partners PrEP	4758 sd ♀/♂ couples	98%	97%	82%
Fem-PrEP	2120 ♀	95%	85%	<40%
VOICE	5029 ♀			
• TDF		90%	87%	30%
• Truvada		91%	92%	29%
• TFV gel		91%	86%	25%

Ambia (review) 2013; Baeten (review) 2013; van der Straten 2012; Baeten CROI 2013; Marrazzo CROI 2013

Measuring Adherence in 017





Timelines

- Protocol development:
 - Protocol development meeting 09-28-11
 - Community consultations
 - Cape Town Oct 2011
 - Pittsburgh Dec 2011
 - Bangkok and Chiang Mai Jan 2012
 - Boston Mar 2012
 - Lima Mar 2012
 - PSRC 04-03-12
 - Version 1.0 07-13-12

Site Activation

□ Clinical Trial Agreement: 07-11-13

□ Sites:

■ Fenway	09-06-13
■ SFDH	09-24-13
■ Pittsburgh*	09-27-13
■ San Juan	11-15-13
■ Chiang Mai	01-06-14
■ Lima	01-15-14
■ Cape Town	02-28-14
■ Bangkok*	06-27-14

*Mucosal immunology subset

017 Activity as of 10-17-14

Site (n)	Screened	Screen Fail	In Screening	Enrolled
Fenway (6)	11	4	0	7*
SFDH (36)	90	52	6	32
Pittsburgh (36)	42	12	1	29
San Juan (6)	13	4	0	7*
Chiang Mai (30)	55	25	2	28
Lima (36)	53	17	2	35
Cape Town (18)	46	28	2	16
Bangkok (24)	32	8	3	22
TOTAL 192	342	150	16	176

* Including replacement ppt/s

Anticipated Timeline

Topic	Date of completion
Accrual	December 2014
Follow up (27 weeks)	August 2015
Data clean up	October 2015
Data lock	November 2015
Primary results	December 2015

Acknowledgements

- MTN is funded by NIAID (5U01AI068633), NICHD and NIMH, all of the U.S. National Institutes of Health
- Gilead Sciences
- CONRAD
- MTN-017 participants

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Lama, Jorge Sánchez, Rosa Infante, Aldo Lucchetti, Javier Salvatierra, Jorge Vergara, Esmelda Montalban, José Gonzales, Eduardo Sánchez, Manuel Villaran, Fanny Garcia, Jessica Rios, Karen Villanueva, Karina Pareja, Monica Sánchez, Carla Porcile, Carmen Sánchez, Richard Teran, Cecilia Correa, Roberto Facho, Peter Brandes, Eduardo Ruiz, Martín Lacherre, Bertha Talaverano, Eliana Díaz, Carolina Moran, Diana Durand, Silvana Torres, Alberto Rondan, Alejandra Flores, Martín Patiño, Esmellin Perez, Robert De la Grecca, Carmela Ganoza, Lily Ganaha, Cecilia Chang, Ricardo Alfaro, Jesus Jurupe, Maria Suarez, Giovanna Solis, Carmen Salinas, Janet Soto, Ronny Tirado, Sonia Minaya, Gustavo Quispe, Roberto Alcantara, Patricia Segura, Medalith Sulca, Yolanda Vidal, Noelia Niño, Luis Castro, Rafael Rosas, Gonzalo Meneses, Daniel Alva, Christian Keller, David Amiel, Julio Dextre, Hector Salvatierra, Martin Patino, Lourdes Cruzado



Thank You!