

HIV ascertainment through repeat home-based testing in the context of a Treatment as Prevention trial (ANRS 12249 TasP) in rural South Africa

Joseph Larmarange, Joanna Orne-Gliemann, Éric Balestre, Collins Iwuji, Nonhlanhla Okesola, Marie-Louise Newell, François Dabis, France Lert

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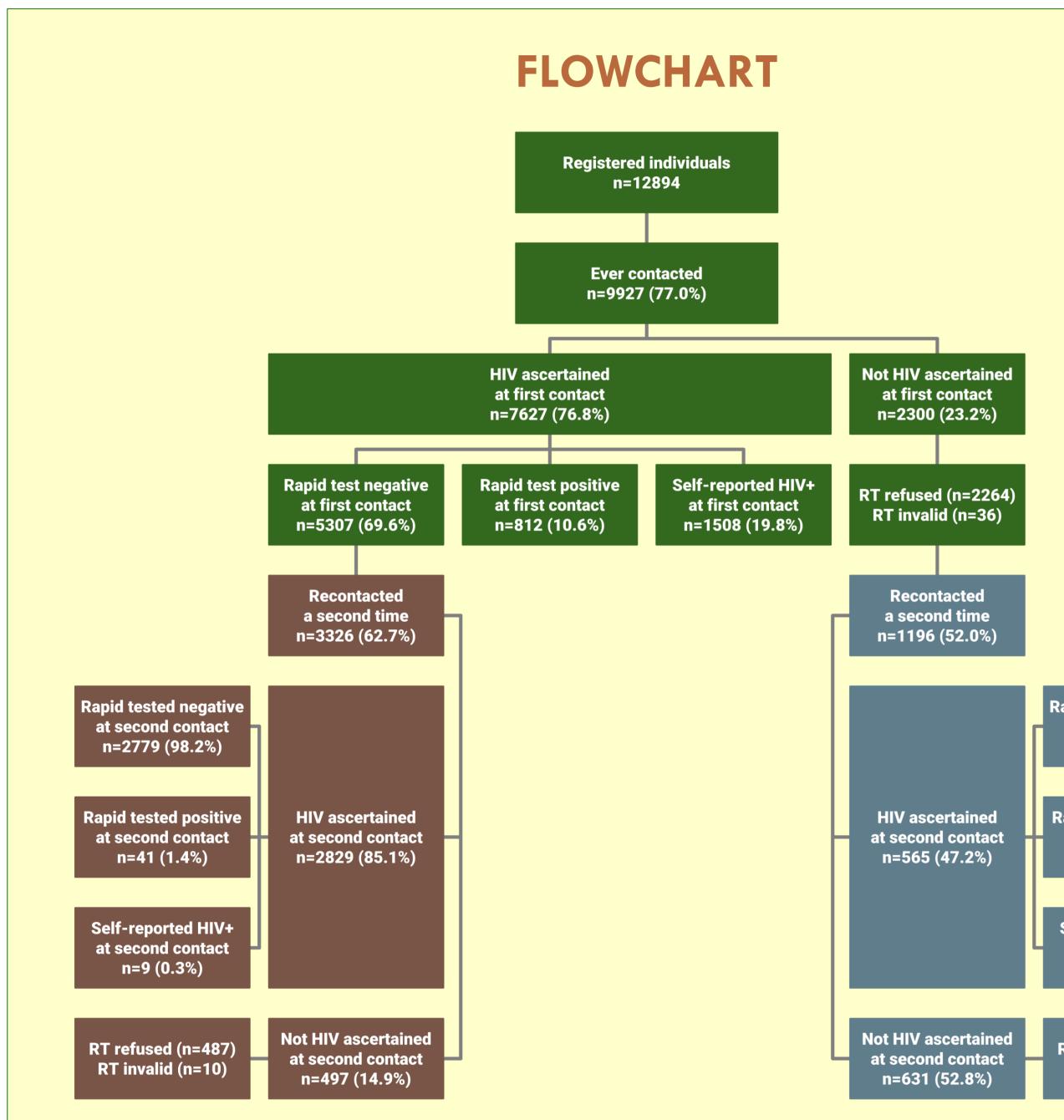
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Antiretroviral Treatment as Prevention • ANRS 12249 Ukuphila kwami, ukuphila kwethu (my health for our health)



Among the 12,894 registered individuals, 9,927 (77.0%, 95% CI: 72.6% - 80.9%) have been contacted at least once; of these, 76.8% (95% CI: 73.0% - 80.3%) have been HIV ascertained at first contact.

A second contact creates an opportunity to find individuals not ascertained at first contact, the 'ever ascertained rate' increased from 76.8% after one contact to 89.2% after two contacts (+12.4, 5237/5868, 95% Cl: 86.7% - 91.3%).

Among individuals tested negative at first contact, repeat ascertainment rate at second contact was 85.1% (95% CI: 82.2% -87.5%).









HIV ASCERTAINMENT THROUGH REPEAT HOME-BASED TESTING IN THE CONTEXT OF A TREATMENT-AS-PREVENTION TRIAL (ANRS 12249 TASP) IN RURAL SOUTH AFRICA

Joseph Larmarange^{1,2*} Joanna Orne-Gliemann^{3*} Éric Balestre³ Collins Iwuji¹ Nonhlanhla Okesola¹ Marie-Louise Newell⁴ François Dabis³ France Lert⁵ for the TasP ANRS 12249 Study Group

1 Africa Centre for Health and Population Studies, University of KwaZulu-Natal, South Africa. 2 Centre Population et Développement (CEPED UMR 196 Université Paris Descartes Ined IRD), France. 3 INSERM / University of Bordeaux, ISPED, Centre Inserm U897- Épidemiologie-Biostatistique, France. 4 Faculty of Medicine, University of Southampton, United Kingdom. 5 Centre de recherche en Épidémiologie et Santé des Populations (CESP, Inserm Unité 1018), Villejuif, France.

CONTEXT

The ANRS 12249 TasP trial evaluates whether HIV testing of all members of a community, followed by immediate antiretroviral treatment (ART) for infected people, will prevent onward sexual transmission and reduce HIV incidence at population level. Ascertaining the HIV status of a high proportion of the population regularly and repeatedly is key to the success of any universal test and treat strategy, as the first step of the HIV cascade.

HIV ascertainment, defined as having taken a rapid HIV test (invalidate and indeterminate results excluded) or self-reporting being HIV-positive, is a process indicator measuring the capacity of the trial to identify the HIV status of the participants and, if required, to refer them to a dedicated trial clinic.

A cluster-randomised trial is being implemented using a phased-approach in the Hlabisa sub-district (KwaZulu Natal, South Africa) where more than 20% of adults are living with HIV. The trial started in March 2012 and is implemented in 10 clusters (5×2) for the first phase to assess the feasibility and acceptability of the test and treat strategies. The HIV testing strategy consists of the implementation of 6-monthly rounds of home-based HIV counselling and testing by dedicated counsellors.

Between March 2012 and March 2014, we have implemented three rounds of home-based HIV counselling and testing in four local communities (clusters) and two rounds in six additional clusters (opened in 2013). At each home visit, individual questionnaires are administered to each consenting participant aged 16 years old or more, who is also offered rapid HIV test.

We report on preliminary results of HIV ascertainment rates at first and second contact, and individual and household-level associated factors.

DISCUSSION / CONCLUSION

Acceptance of regular and frequent HIV testing is key to the community-based efficacy of treatment-as-prevention initiatives (universal test and treat) in settings with very high incidence.

Our preliminary data show that, as long as individuals are successfully re-contacted, acceptability of repeat home-based HIV testing is high (85%) among individuals previously tested negative at home. It is important to stress that only two to three rounds of home-based HIV testing have been implemented to date and it is possible that a participation fatigue will emerge over time.

Our factors analysis show that home-based testing may not capture and enroll young and active populations, especially men. Lower chance of HIV ascertainment among adults aged 20 to 40 years and among the higher educated suggests that this strategy might be less acceptable among people who can possibly access HIV testing elsewhere.

Although statistical significance of the associations at second contact is less, patterns looks similar between ascertainment at first contact and at second contact among individuals not ascertained at first contact.

Differences between those ascertained and not ascertained at second contact among those tested negative at first contact are minimized: one of the keys to engaging individuals on a repeat HIV testing pathway within the trial remains the acceptability of the first home-based HIV test.

The effect of risky sexual behaviour seems to depend on sex: among men, a more self-reported risky behaviour is associated with increased test refusal, while the opposite is observed among women. Additional analyses are required to improve our understanding of the sexual activity and behaviour of these individuals who declined home-based HIV testing despite several attempts at contacting them and were thus never ascertained. This will help assess to what extent repeated home-based HIV testing campaigns could be an efficient strategy to reach the populations most at risk of transmitting HIV, in particular mobile individuals.

Repeat home-based HIV testing appears as one of the effective strategies to achieve a high level of testing coverage in these high prevalence rural areas. However, combined HIV testing approaches could be required to achieve universal testing in socially heterogeneous communities.

TASP STUDY GROUP (alphabetical order): Till Barnighausen, Sylvie Boyer, Alexandra Calmy, Marie-Laure Chaix, Francois Dabis (co-PI), Rosemary Dray-Spira, Ken Freedberg, John Imrie, Collins Iwuji (South Trial Coordinator/Trial Physician), Sophie Karcher, Joseph Larmarange, France Lert, Richard Lessells, Kevi Naidu, Colin Newell, Marie-Louise Newell (co-PI), Nonhlanhla Okesola, Tulio de Oliviera, Joanna Orne-Gliemann (North Trial Coordinator), Deenan Pillay (co-PI), Bruno Spire, Frank Tanser, Rodolphe Thiebaut, Johannes Viljoen, Thembelihle Zuma.

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Rapid tested negative at second contac n=393 (69.6%)

Rapid tested positive nt second contact n=79 (14.0%)

Self-reported HIV+ at second contac n=93 (16.5%)

RT refused (n=629 RT invalid (n=2)

Sex ് men ♀ women Age at 1st contac 15-19 20-29 30-39 40-49 50-59 60 + missing **Educational level** completed secondary some secondary primary or less / don³ missing Household's assets low middle high missing HIV test before first of < 6 months 6 to 12 months > 12 months never tested missing / refused Sexual partner(s) las 2 sexual partners or r 1 sexual partner† none / never had sex missing / refused / de

Condom use at last i yes no †

never had sex

missing †

Knowing someone F yes missing









FACTORS ASSOCIATED WITH HIV ASCERTAINMENT

	Ascertainment at 1 st contact			Ascertainment at 2 nd contact among tested negative at first contact (n=3,326)			Ascertainment at 2 nd contact among not ascertained at first contact (n=1,196)		
	(n=9,927)								
	%	aOR	95% Cl	%	aOR	95% CI	%	aOR	95% C
	***	**		**	***		-		
	74.1	1		82.3	1		46.6		
	78.1	2.52	1.54-4.11	86.2	1.29	1.05-1.59	47.5		
	***	***		**	***		***	**	
	78.5	1		81.7	1		53.8	1	
	70.4	0.66	0.55-0.79	81.2	0.94	0.70-1.26	37.7	0.63	0.41-0.98
	77.0	0.77	0.62-0.95	83.9	1.12	0.74-1.68	54.6	1.16	0.70-1.9
	85.1	0.95	0.74-1.23	87.1	1.44	0.98-2.11	48.3	0.61	0.34-1.10
	85.4	0.89	0.68-1.18	87.5	1.49	1.05-2.11	50.0	0.56	0.29-1.0
	86.3	0.98	0.75-1.30	87.7	1.55	1.15-2.08	61.7	0.96	0.52-1.78
	39.2	0.57	0.41-0.79	90.6	2.24	0.86-5.83	23.4	0.52	0.20-1.37
	***	***		**			***	***	
/ higher	71.8	1		81.5			37.6	1	
, -	78.5	1.32	1.15-1.50	83.2			54.1	1.89	1.33-2.69
t know	87.6	2.23	1.87-2.67	87.4			61.0	2.81	1.72-4.59
	9.8	1.55	0.75-3.22	77.8			4.2	0.87	0.14-5.54
ore category	-			-			*		
. ,	77.2			85.7			51.3		
	77.0			84.4			47.8		
	75.8			85.1			38.9		
	86.8			70.0			25.0		
ontact ‡	***	***							
÷	68.4	1							
	85.3	2.57	2.18-3.03						
	86.0	2.54	2.22-2.90						
	81.2	1.67	1.42-1.96						
	19.1	1.69	1.05-2.71						
12 months	***	**		_			***		
ore	75.5	1		81.9			43.1		
	80.7	י 1.34 ס'	1.08-1.67	85.7			49.4		
	00./	♀ 0.55	0.35-0.85	00.7					
	82.0	° 2.25	1.38-3.68	85.5			52.8		
	02.0	♀ 0.39	0.24-0.63	0010					
on't know	44.9	d [™] 0.93	0.58-1.50	80.6			30.4		
		♀ 0.35	0.21-0.59						
itercourse	***	**		_			***		
	76.8	1		82.8			43.8		
	82.1	ਾ ਹੋ 0.97	0.75-1.25	86.1			52.3		
		♀ 1.08	0.92-1.26						
	78.0	of 3.21	1.74-5.91	83.0			49.1		
		Q 1.53	1.10-2.14						
	42.1	o [*] 0.74	0.37-1.49	83.0			26.3		
		Q 1.08	0.77-1.52						
V+ in family	***	***		*			***	*	
	79.1	1		84.3			49.8	1	
	83.4	1.29	1.14-1.46	87.3			50.4	1.04	0.77-1.40
	1.8	0.01	0.01-0.02				1.5	0.03	0.00-0.43

%: HIV ascertainment rate within modality (univariable analysis). aOR: adjusted odd ratio (multivariable analysis). Cl: confidence interval (odd ratio). -p > 0.05; * 0.01 $; ** 0.001 <math>; *** <math>p \le 0.001$ (Chi² test for univariable; Wald or log likelihood test for multivariable) *†* stratified by sex (multivariable analysis); *‡* variable used only for first contact analysis.

Only variables with p≤0.20 in univariable analysis have been included in multivariable logistic models using a step-by-step backwards method (Wald criterion). Interactions between sex and others risk factors were tested and potential confounding were checked. Clusters effect was taken into account by using random effects models (except for first model who did not converge).