## Point-of-care Testing for Chlamydia Infection Data extraction – Diagnostic accuracy

Reviewer ID:	Date:
Administration details for study:	
Study ID:	Study design:
Multicentre study:  Yes. Number of centres  No	☐ - RCT  All patients randomised to index vs comparator. All also receive the reference standard
Country/countries:	Direct (head to head) comparison. All patients receive index test, comparator and reference standard
<b>Setting</b> (e.g. primary care, GUM clinic, community health point):	Duration of study:
Funding details government / private / manufacturer / other (specify):	Study start/end dates:
Additional info:	Length of follow-up:
Aim of study:	

Comparisons:			
Chlamydia Rapid test vs other POCT comparator and reference standard			
Specify index test			
Specify comparator			
Specify reference standard used			
Chlamydia Rapid test vs NAAT comparator (i.e. the reference standard)			
Specify index test			
Specify reference standard used			
Outcomes reported:			
Test performance results	Data provided for relevant subgroup:		
Acceptability of the test to patients/healthcare staff (delete as appropriate)	☐ Those aged <25 years old ☐ Men who have sex with men (MSM) ☐ Sex workers		
☐ Interpretability of the test	High-risk African populations		
Inclusion criteria:			

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EXC	1151011	criteria:

Characteristics of participants:					
	Index test	Comparator 1	Comparator 2	All	
Enrolled					
Received tests					
Received reference					
standard					
Analysed					
Number of					
uninterpretable					
tests					
Lost to follow-up					
Age					
(mean/median,					
SD/IQR range)					
Sex	F:	F:	F:	F:	
	M:	M:	M:	M:	
% aged < 25					
% MSM					
% sex workers					
% other high risk					
groups (please					
specify)					

Additional information on participants:

Characteristics of the tests:			
Index test: Chlamydia Rapid Test	Manufacturer, country:		
Number of samples taken per patient:	Time taken till test result available:		
Sample(s) obtained by:	Setting where sample was collected:		
☐ Voided urine	General practitioner		
Urethral swab	GUM clinic		
Endocervical swab	Family planning centre		
☐ Vaginal swab (self taken)	Acute care		
☐ Vaginal swab (taken by practitioner)	Other community health point. Please specify (e.g		
Other (specify)	pharmacy, youth club)		
Positive test result defined as:  Additional information on test:			

Comparator test 1:	Manufacturer, country:
Type of test:	Time taken till test result available:
POCT	
NAAT	
Other (please specify)	
Number of samples taken per patient:	
Sample(s) obtained by:	Setting where sample was collected:
Voided urine	General practitioner
Urethral swab	GUM clinic
Endocervical swab	Family planning centre
☐ Vaginal swab (self taken)	Acute care
☐ Vaginal swab (taken by practitioner)	Other community health point. Please specify (e.g.
Other (specify)	pharmacy, youth club):
Positive test result defined as:	
Additional information on test:	

Characteristics of the tests:	
Comparator test 2:	Manufacturer, country:
Type of test:	Time taken till test result available:
POCT	
NAAT	
Other (please specify)	
Number of samples taken per patient:	
Sample(s) obtained by:	Setting where sample was collected:
☐ Voided urine	General practitioner
Urethral swab	GUM clinic
Endocervical swab	Family planning centre
☐ Vaginal swab (self taken)	Acute care
☐ Vaginal swab (taken by practitioner)	Other community health point. Please specify (e.g
Other (specify)	pharmacy, youth club):
Positive test result defined as:	
Additional information on test:	
Reference standard test:	Manufacturer, country:
Type of NAAT:	Time taken till test result available:
Polymerase chain reaction (PCR)	
Ligase chain reaction (LCR)	Time interval between index test and reference standard
Strand displacement amplification	test:
(SDA)	
Transcription mediated amplification (TMA)	

Sample(s) obtained by:	Setting where san	nple was collected:	
☐ Voided urine	General practi	tioner	
Urethral swab	GUM clinic		
Endocervical swab	Family plannii	ng centre	
☐ Vaginal swab (self taken)	Acute care		
☐ Vaginal swab (taken by practitioner)	Other commun	nity health point. Ple	ease specify (e.g
Other (specify)	pharmacy, youth	club)	
Positive test result defined as:			
Additional information on test:			
Results:			
Index test	Comparator 1	Comparator 2	Reference
			standard
N in analysis			
True-positives			
False-positives			
True-negatives			
False-negatives			

Number of samples taken per patient:

Sensitivity

Specificity				
Positive LR				
Negative LR				
PPV				
NPV				
Diagnostic odds				
Additional informa	ation on results (e	e.g. contradictory result	s resolved by):	
Adverse events:				
General information	on on adverse eve	nts:		
Adverse events	Index test	Comparator 1	Comparator 2	Reference standard

Acceptability and interpretability of tests:	
Additional study information:	