

# HIV Prevention England Conference 2017

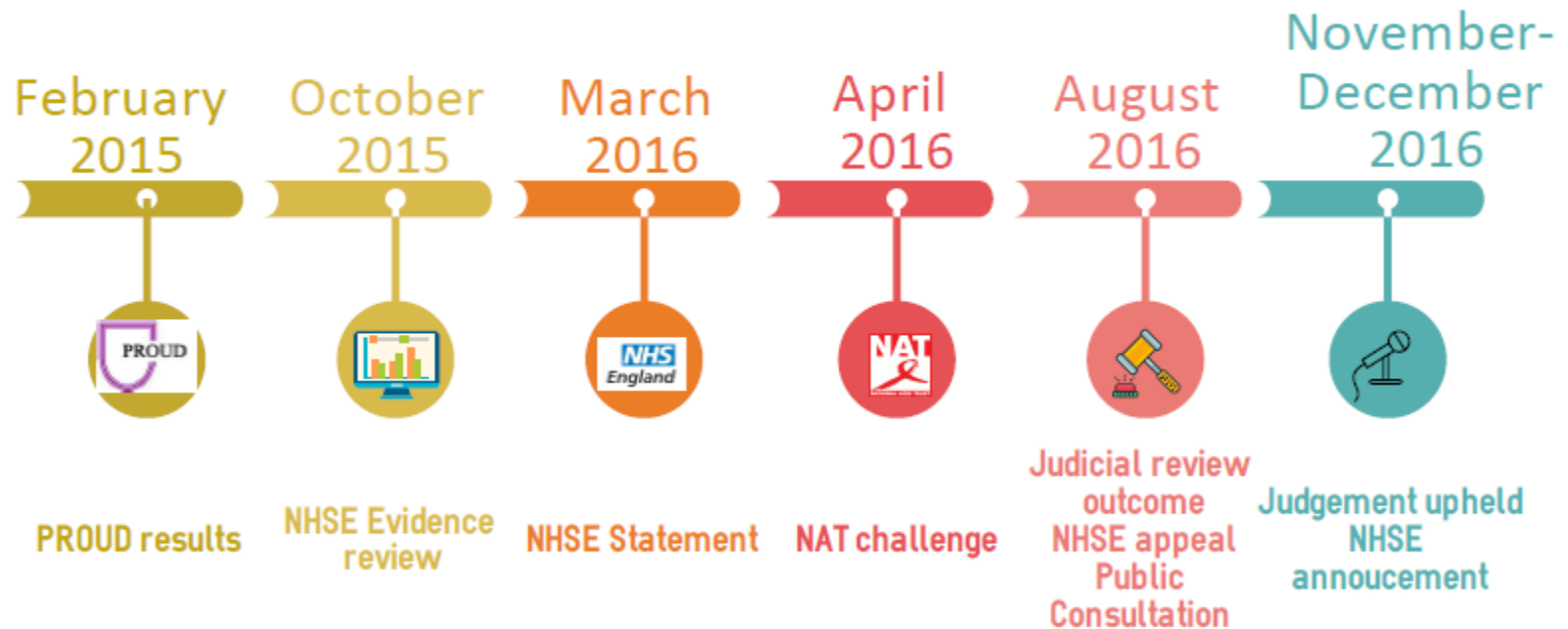
## Update on the PrEP Impact Trial

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# UK PrEP timeline



<https://www.england.nhs.uk/2016/11/update-on-prep/>  
<https://www.england.nhs.uk/2016/12/hiv-prevention-programme/>

# PrEP IMPACT Trial



£10m



10,000



3 years



## Address key outstanding questions

- 1) Proportion eligible for PrEP?
- 2) How to identify, engage and maintain all people eligible for PrEP?
- 3) Proportion accepting offer of PrEP?
- 4) Proportions choosing daily or intermittent dosing?
- 5) Duration of PrEP use?
- 6) Impact on HIV incidence?
- 7) Impact on STI incidence?

### Primary objective

To measure PrEP-eligibility, PrEP-uptake, duration of PrEP-eligibility and duration of PrEP-use among Genitourinary Medicine (GUM) clinic attendees

# Inclusion criteria

## **Key principles of the inclusion criteria:**

Include all persons at high risk of HIV:

1. Higher risk sexual behaviour
2. HIV positive partner
3. Partner of unknown status and at high risk of HIV

# Inclusion Criteria

## 1 Cis- and transgender MSM and trans women

- a) HIV negative test in previous year
- b) Report condomless sex in the previous 3 months
- c) Affirm likelihood of CSI in the next 3 months

## 2 HIV negative partner of an HIV positive person

- a) HIV positive partner not known to be virally suppressed
- b) CSI anticipated before treatment of HIV positive partner takes effect

## 3 HIV negative person

Clinically assessed and considered to be at similar risk of HIV acquisition as those with a serodiscordant partner who is not known to be virally suppressed

## First visit

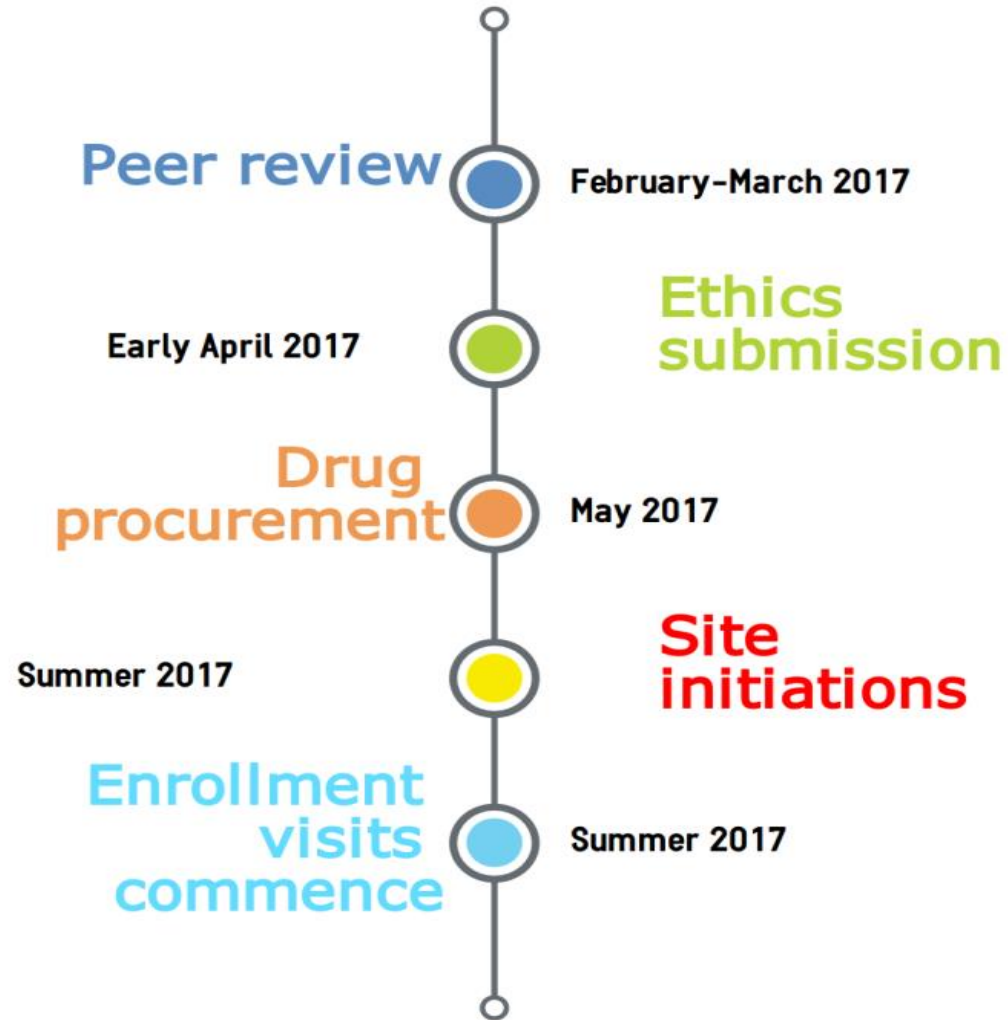
- Baseline tests (HIV test, STI tests, Kidney function)
- 3-month PrEP prescription (daily or EBD)
- Routine data collection



## Follow up visits (1 month after starting and every 3 months)

- HIV/STI tests according to routine practice
- Hepatitis C test according to routine practice
- Kidney function
- Combination prevention
- Safety check
- PrEP prescription (daily or EBD)
- Routine data collection

# Proposed trial timeline



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Address key outstanding questions



Based on proposed eligibility criteria



Mitigate against future  
geographical inequity of access



Support regular clinical risk  
assessments as part of risk reduction



Use GUMCAD as data collection spine



# PrEP Impact Trial



and



Public Health  
England

Protecting and improving the nation's health

**Chief-Investigator:** Professor Brian Gazzard – Chelsea and Westminster NHS Foundation Trust

**Co-Investigators:** Dr Anne Sullivan, Professor Sheena McCormack, Professor Noel Gill, Dr Monica Desai, Dr John Saunders, Dr Valerie Delpech

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