1 Title: Acceptability, facilitators and barriers to point of care HIV testing in a homeless-focused 2 service in Gloucestershire: A qualitative evaluation 3 4 **Short title:** Point of care HIV test: facilitators and barriers 5 6 Authors: E Corker¹, F Lorencatto¹, N Anderson¹, M Gobin², S Scott³, S Michie¹ & G Angel² 7 8 **Author affiliations:** 9 1. Centre for Behaviour Change Dept of Clinical, Educational, and Health Psychology, University 10 College London, 1-19 Torrington Place, London WC1E 7HB, UK. 11 2. Public Health England, Wellington House, 133-155 Waterloo Rd, London SE1 8UG, England. 12 3. Gloucestershire County Council, Shire Hall, Westgate Street, Gloucester, GL1 2TG. 13 14 Corresponding author: Elizabeth Corker, email: e.corker@ucl.ac.uk Postal address: Room 353, 1-19 15 Torrington Place, Department of Clinical, Educational & Health Psychology University College London 16 London, WC1E 6BT. 17 18 **Abstract** 19 20 Late HIV diagnosis increases risk of onwards transmission, morbidity and mortality. Rapid Point of 21 Care Testing (POCT) reaches people never tested and people living with HIV who are undiagnosed. A 22 pilot study introduced HIV-POCT to one service in Gloucestershire, England. This study explored 23 acceptability and feasibility of HIV-POCT in this context, from the perspective of service providers 24 and users. Eleven semi-structured interviews with service users and a focus group with three service 25 providers were conducted. The Theoretical Framework of Acceptability and the Theoretical Domains 26 Framework were used to design the topic guide and analysis. Acceptability of HIV-POCT was high. 27 Seven facilitators were identified (e.g. understanding test purpose and process), alongside two

- 1 potential barriers, one relevant to service providers and users (anxiety) the other in relation to
- 2 service users (stigma). To maximise implementation of HIV-POCT, healthcare providers require
- 3 appropriate training and supervision to offer and administer POCT.

Main text

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1. Introduction

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5 Undiagnosed and late diagnosis of HIV is associated with negative outcomes for onwards 6 transmission, morbidity and mortality (1-4), and has significant implications for public health. Public 7 Health England (PHE) define late HIV diagnosis as "a CD4 cell count <350 cells/mm3 within 91 days 8 of... HIV diagnosis" (page 6, Public Health England, 2019). Rates of late HIV diagnosis are high (49%) 9 in Europe (6) with the proportion of late diagnoses in England being 43% (5), with an estimated 10 6,700 people living with undiagnosed HIV (5). 11 12 Risk of contracting HIV is higher for people who are homeless, or use intravenous drugs (7–9). Being 13 diagnosed within 1 year of HIV infection (10) helps reduce rates of HIV infection (3,8,11) and testing 14 is available in numerous settings (e.g. sexual health clinics, charities, GP surgeries, pharmacies) (12). 15 16 Literature reviews suggest barriers for testing within the general population exist at: service user 17 (e.g. low risk perception, fear of positive results, negative consequences, low awareness of services, 18 low HIV knowledge, perception of cultural appropriateness or quality of test, and low perceived 19 knowledge of HIV in clinical staff (13,14)); service provider (e.g. reluctance to address HIV and offer 20 test; lack of confidence or skill in testing; lack of awareness of HIV epidemiology and feeling 21 uncomfortable with post-test counselling (13,14)) and institutional or policy levels (e.g. financial and 22 human resources (13)). Additionally, for people who are previous or current injecting drug users, 23 standard HIV testing can be challenging due to difficulties in finding a viable vein (15,16). 24 25 Evidence suggests that rapid point of care testing (POCT) (a finger prick test not requiring samples be

sent to a laboratory and can be performed where or near a patient presents) (17) reaches people

who have never been tested and people living with undiagnosed HIV (18). An evaluation in North West England indicated that targeting marginalised groups through community based POCT was feasible and acceptable to service providers and users and successfully reached groups that would not have otherwise been tested (19). Community POCT is recommended to overcome barriers such as stigma at using hospital-based services (British HIV Association, 2008), although concern has been expressed that rapid testing could lead to rapid counselling and testing without consent (11,20). One area in the South West of England (Gloucestershire) has seen a number of successive periods (3 year rolling averages) of increases in the proportion of late diagnosis, resulting in 65% of HIV diagnosis defined as late, statistically significantly higher than the England average (21). Descriptive epidemiology identified risk factors potentially associated with late diagnosis, including people who inject drugs. A cohort of known cases of HIV, who have risk factors associated with poor treatment adherence and onward infection has been identified and all but one of these are users of local homeless healthcare services. A pilot study introduced HIV-POCT to a homeless-focussed service in Gloucestershire as one way to reduce the rate of late HIV diagnosis in this population. Implementation of new healthcare interventions is rarely a straight-forward or linear process (22). Investigating acceptability, barriers and facilitators to implementation in a pilot study can identify challenges to be addressed to improve acceptability, implementation and uptake before wider rollout, in line with Medical Research Council complex intervention guidance (23). Previous research has differentiated barriers and facilitators by levels in the system: individual/groups/organisations (13,14), providing valuable insights. However, as health behaviour interventions are delivered as part of complex systems, it may be helpful to conceptualise barriers and facilitators in line with ecological systems theory: individual/micro-systems (service users and providers), exosystems/service level (service-wide issues and communication between systems), and macro-

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systems/population levels (socio-political landscape) (24), to understand how systems inter-relate in

the context of healthcare interventions.

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4 Frameworks in the behavioural and implementation sciences facilitate exploration of acceptability

and barriers and facilitators to implementation of healthcare interventions. The Theoretical

Framework of Acceptability (TFA) (25), proposes seven different dimensions of acceptability (e.g.

ethicality, burden, intervention coherence), and can help inform more comprehensive, multi-

dimensional assessments of prospective, concurrent, and/or retrospective acceptability of

healthcare interventions. The Theoretical Domains Framework (TDF) (26) synthesises 33 theories of

behaviour change into 14 domains representing individual, socio-cultural and environmental barriers

and enablers to implementation and behaviour change. Both frameworks have been applied to

explore barriers/enablers and acceptability of a wide range of healthcare interventions (TFA (27,28)

13 and TDF (29–32)).

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This pilot study aims to apply the TFA and TDF to investigate acceptability of barriers and facilitators

to implementation of HIV-POCT from the perspective of service users and healthcare providers at

individual/micro-system, exo-system, and macro-system/population levels.

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2. Methods

Design: Service users participated in a one-to-one semi-structured qualitative interview and

healthcare providers took part in a focus group. Service users were offered a one-to-one interview

to ensure that a suitable time and date could be found for each participant and as an

acknowledgement that the interview may cover confidential information that service users would

feel uncomfortable sharing in a focus group. A focus group was offered to healthcare providers to

ensure that discussions were possible regarding the feasibility and acceptability of the POCT

- 1 amongst those administering it. Ethical approval was granted from UCL Research Ethics Committee
- 2 (Ref: 15851/001).

- 4 2.1. Sample
- 5 We had two participant groups: 1) people currently accessing homeless services within
- 6 Gloucestershire, (according to Local Authority figures, 194 households were classified as 'homeless'
- 7 in 2019 (33), along with 198 injecting drug users (Offer, Willis and New, 2015)); 2) healthcare
- 8 providers responsible for delivering POCT (N= 4).
- 9 Over the four-month pilot study period, new or returning service users, over the age of 18, where
- 10 the healthcare provider judged they had adopted one or more risky behaviours which could expose
- them to HIV infection (i.e. injecting drug use or sex working) were offered HIV-POCT.
- 12 Service users offered POCT were approached by healthcare providers during routine visits with
- information sheets and an invitation to participate in the present study. A designated 'go to'
- 14 healthcare provider assisted researchers in identifying healthcare providers responsible for
- delivering POCT for the researcher to contact with an information sheet containing the researcher's
- 16 details.
- 17 Target sample size estimations were based on recommendations for qualitative research and
- principles of thematic data saturation (35). Data saturation (the point where new data does not
- 19 contribute any new information) is deemed more meaningful in nonprobablistic research for
- 20 estimating a sample size, than a power calculation (36). The estimated point of data saturation
- varies between six (36) and 13 interviews (35). Therefore, we set a target of 13 interviews with
- service users. It was not possible to set this target with the focus group, due to the total N (4) of the
- 23 healthcare provider population.
- 24 2.2. Measures

- 1 The following quantitative information was collected from the patient record system, to provide
- 2 context to the qualitative information for the service user group: age; sex; ethnicity; type of
- 3 accommodation; presence of risk factors for HIV infection (bought or sold sex; men who have sex
- 4 with men; sexual contact with someone known to be HIV positive; injecting drug use; prison record);
- 5 acceptance or refusal of HIV-POCT, and test result.
- 6 2.3. Interviews and focus group
- 7 Informed consent was gained for data collection including audio recordings and using the data for
- 8 research purposes. All one-to-one interviews took place in one sitting and were held in a private
- 9 room within the service providing POCT. One focus group was held with healthcare providers. The
- interviews and focus group were conducted by an experienced researcher and transcribed verbatim.
- 11 A £10 remuneration was offered to each service user participant.
- 12 The two-part interview and focus group topic guides were semi—structured and based on the TFA
- and TDF. Part one focused on POCT acceptability, and included at least one question per domain of
- the TFA (25). Part two focused on potential barriers and facilitators of receiving or delivering the
- 15 POCT, with questions structured around the domains of TDF (26). At least one question per domain
- of the TDF was included. To avoid potential repetition, questions were not exclusively assigned to a
- domain, and one question may have been designed to gain information related to conceptually
- 18 similar domains within the TDF and TFA, for example, the same question was designed to be
- 19 relevant for the TFA domain 'affective attitude' and the TDF domain 'emotion'. In total 15 questions
- were included in the topic guides for the one-to-one interviews (not including prompts) and eight
- 21 questions were included in the topic guide for the focus group (not including prompts). Table 1
- includes definitions of the domains from each framework and an example question from the
- 23 interview and focus group that tap into each domain. Topic guides for service users were piloted
- through a Patient and Public Involvement group. For full topic guides see Supplementary file 1.

Table 1 here.

- 3 2.5. Analysis
- 4 Analysis was based on guidance for analysing qualitative data based on the TDF (37), and Thematic
- 5 Analysis (38) using NVivo12. This resulted in a five-stage combined deductive and inductive analysis
- 6 (39) approach:
- One experienced qualitative researcher familiarised themselves with transcripts of the
 interviews and focus group. A codebook was generated based on the TFA and TDF, giving
 examples of codes and associated data (participants quotations).
 - 2. Deductive framework analysis: Participant responses were deductively coded to the domains of the TFA and/or TDF that they were judged to best represent (e.g. 'it's a bit nervewracking' (pt 1) was coded to the TDF domain 'emotion'), by a single researcher, and subsequently reviewed by a second researcher with experience in application of the TFA and TDF. Discrepancies were resolved through discussion (40).
 - 3. Inductive thematic analysis: Similar responses coded to the same domain from each framework were grouped together, and a summary theme label inductively generated.
 - 4. The final list of themes was subjected to member checking by a second researcher to assess whether they: 1) reflected shared meaning of quotes and 2) the theme belonged to the domain it was assigned. Where discrepancies arose, theme labels were refined or reassigned to a different domain until consensus was reached.
 - 5. Themes were inductively analysed in relation to how they described processes belonging to an individual, the service or wider, or reflecting socio-political systems. Themes can exist within different levels, for example, a barrier to acceptance may exist on the individual level, as well as on the population-level system, highlighting the complexity of the impact of healthcare interventions. A map of the themes was produced.

1 2 **Results** 3 4 3.1. HIV point of care testing – uptake and results 5 When the study period ended, 14 people had been identified by the healthcare provider as having 6 adopted one or more risky behaviours which could expose them to HIV infection and were offered 7 POCT. Of these 14 people, 11 (79%) accepted and the POCT were administered by nurses working within the homeless service. All POCT tests undertaken were 'unreactive': indicating no HIV 8 9 infection. 10 11 Eleven service-users participated in the interviews, 10 who had accepted POCT and 1 who refused, 12 see Table 2 for demographic details. One focus group was held with three service providers: two 13 nurses and one health protection practitioner. 14 15 Table 2 here. 16 17 3.2. Themes across both frameworks 18 Table 3 presents themes inductively generated from service users and healthcare providers with 19 supporting exemplary quotes. No themes emerged that were not covered by TFA or TDF domains. 20 Therefore, themes are mapped onto the TFA and TDF domains to which they correspond, and in 21 relation to the TDF categorised as either a barrier or facilitator to delivery and uptake of POCT; in 22 relation to the TFA, a categorisation as a 'facilitator' demonstrates acceptability. A number of 23 common themes were identified between service user interviews and the service providers focus 24 group. Many of the themes that emerged for certain domains of the TFA and TDF were similar (e.g.

knowledge and intervention coherence) thus the decision was made to merge these together and

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present them jointly.

Table 3 here

3.3. Facilitators and acceptability of POCT

Of the 11 themes, seven were facilitators, (see Table 3), enabling or influencing participants uptake of POCT, and contributing to its acceptability. Service users found POCT acceptable partly due to understanding the purpose and process of POCT, believing staff were able to perform POCT and believing POCT would be effective, highlighted through the TDF domains 'knowledge', 'beliefs about capabilities' and 'optimism', along with the TFA domains 'intervention coherence', 'self-efficacy' and 'perceived effectiveness'. Further facilitators were: belief in a responsibility to take a HIV test (social and professional identity (TDF) and ethicality (TFA)), being presented with a choice of taking the test (memory, attention and decision making processes (TDF)) and not having to carry out additional behaviours, such as going to a scheduled blood test appointment (environmental and contextual resources (TDF) and burden (TFA)).

With regards to the healthcare providers, facilitators of delivering POCT included understanding the purpose of the test and believing they were able to administer it, evidenced by themes related to knowledge (TDF) and intervention coherence (TFA), as well as themes related to beliefs about capabilities (TDF) and self-efficacy (TFA). Further facilitators included: believing that POCT would be effective (optimism (TDF) and belief about efficacy (TFA)); belief that POCT would impact commissioning (beliefs about consequences (TDF)); desire to provide a good service, arising from contact with this population and understanding the positioning of their clinical role, (social and professional identity (TDF) and ethicality (TFA)); ability to fit the test in with other clinical duties (environmental context (TDF) and burden (TFA)) and abilities to make clinical judgements regarding who would be tested (memory attention and decision making processes (TDF)). These results suggest the healthcare provider sample found POCT to be acceptable and feasible to deliver.

1 2 3.4. Barriers 3 4 As only two barriers to uptake of POCT were found, this summary also includes the two themes that 5 were categorised as both facilitators and barriers (mixed). 6 7 For the service user group, a potential barrier to accepting POCT was awareness of negative 8 attitudes towards people living with HIV (emotion (TDF) and affective attitude (TFA)). Mixed (both 9 facilitator and potential barrier) themes related to service user's preference to retain agency over 10 receiving POCT (memory, attention and decision-making processes (TDF)) and uncertainty regarding 11 what would happen in the event of a positive result (emotion (TDF) and affective attitude (TFA)). 12 13 With regards to the healthcare providers, uncertainty regarding what would happen in the event of 14 a positive result (emotion (TDF) and affective attitude (TFA)) was a potential barrier, and agency 15 around using clinical judgement to offer the test was both a facilitator and potential barrier if this 16 agency were to be removed (memory, attention and decision making processes (TDF)). 17 18 3.5. Systems 19 Themes were differentiated according to the system they related to (24)) to demonstrate how they 20 operate across the different levels (Figure 1). The majority of themes sat under the micro-21 system/individual level and demonstrated that, potentially action at this level could determine 22 acceptability of POCT for service users. Service level themes were relevant to the healthcare 23

providers which may represent further facilitation, or barriers to uptake and acceptability. One

theme, 'stigma' was categorised at the micro-system/individual level and the macro-

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system/population level, demonstrating how this operates at both levels.

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1 2 Figure 1 here 3 4 **Discussion** 5 6 This qualitative study identified that HIV-POCT is acceptable to both service users and healthcare 7 providers in Gloucestershire. Facilitators to service implementation and uptake included 8 understanding the test, the processes, ease and convenience of testing and perceived effectiveness 9 of the test. Two barriers were identified: stigma and anxiety. These findings support feasibility and 10 scalability of HIV-POCT for high-risk, priority population groups, such as people who inject drugs or 11 are homeless (15,16,18). 12 13 Previous research suggested barriers to testing such as: low risk perception, fear of positive results and negative consequences (13,14) within service user groups. Similar concerns were found in the 14 15 present study; however, they did not lead to refusal of POCT, potentially as the facilitating factors 16 held more weight in the decision process. Previous research found barriers among healthcare 17 providers to be a lack of confidence or skill in testing and discomfort with post-test counselling 18 (13,14). Similar themes were found within the present study; however, they did not impede the 19 delivery of POCT. Additionally, potential practical barriers with tests requiring a blood sample due to 20 difficulties in finding a viable vein, an issue for this sample as the majority were current or former 21 injecting drug users (15,16), are overcome with POCT. 22 23

Concerns among service users were raised regarding awareness of negative attitudes held within the community towards both people living with, and people being tested for, HIV. This illustrates how stigma can influence HIV testing and therefore, transmission (11,41–43). Another potential concern within this sample, also reflected in the literature is the risk of 'testing without consent' (11,20),

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1 providing POCT as a choice was important for this sample; it was suggested that integrating POCT

into routine testing may decrease uptake, and damage relationships with clinical staff.

4 For healthcare providers potential barriers centred on uncertainty of the process in the event of a

reactive test result. A protocol for receiving a reactive test was set out, however, it was not tested as

part of this study, as no reactive tests were administered, and healthcare providers expressed

concern regarding availability of onward services. In contrast, service users felt reassured and

confident in healthcare providers abilities to facilitate any further interventions, which was

important in managing feelings of anxiety.

Conceptualisation of the themes on a systems level allows for an understanding of how the study impacted on different (albeit related) sectors of the whole system. Additionally, there is concern that using frameworks to analyse qualitative data may miss important information regarding relationships between themes (39); the ecological systems analysis gave an insight into relationships between themes. Given the scope of this study, it is unsurprising that the majority of themes fall within the micro-system/individual level. However, it is important to understand how actions taken regarding HIV-POCT at an individual level can impact on levels further up the system in the form of transmission rates and stigmatising attitudes. Therefore, messages and interventions designed to impact on an individual level need to be consistent with aims and messages on a macro level, such

4.1. Policy and research implications

as decreasing rates of HIV transmission and stigmatising attitudes.

Results from this study suggest that introducing POCT into local services serving populations at risk of contracting HIV would be acceptable for healthcare providers and service users, with the following caveats. It is important that POCT is presented as a choice and that clinical staff have the skills to understand and administer the test, thereby managing anxiety. Clinical staff also need to be

1 aware of the potentially stigmatising impact of being offered POCT and should ensure that 2 confidentiality is maintained, and communication handled in a sensitive manner. 3 4 Further understanding is needed for healthcare providers about the onward clinical journey of a 5 service-user in in the event of a reactive test. Understanding could come in the form of training, 6 preceded by qualitative research to understand barriers and facilitators. 7 8 Additionally, research is required to better understand the impact of any interventions implemented 9 in the micro-system/individual level on levels further up the ecological system. A mixed methods 10 approach could be used to quantitatively track uptake of POCT and HIV transmission rates, 11 combined with a qualitative analysis of attitudes towards HIV within society. 12 13 4.2. Limitations 14 The total service user group sample of 11 did not meet our target of conducting 13 interviews with 15 service users. Additionally, only one participant in the service user group refused POCT, due to 16 having recently been tested for HIV. Although the high uptake of POCT may reflect high levels of 17 acceptability within this sample, this may have reflected issues with data saturation as it made 18 identifying barriers difficult, as none were cited as reasons not to take up the offer of POCT. These 19 barriers could more accurately be described as potential barriers, and services could benefit from 20 ensuring that they are attended to before implementing POCT. 21 22 Although the overall number of people offered POCT within the study was small, the data suggest 23 that it targeted people considered to be at high risk of HIV infection and late diagnosis i.e. people

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who inject drugs and those who are homeless.

1 A limitation of the data collection procedure was the omission of recording if there was a pre-

2 existing relationship between the individual service user being offered the POCT and the healthcare

3 provider administering the POCT. Collection of this data may have offered an insight into who might

be best placed to deliver HIV-POCT and we recommend this is considered in future research.

Additionally, we did not have data relating to if service users in this sample had access to POCT for

blood borne viruses such as Hepatitis C (HCV). Collecting data on availability and uptake of other

POCT may aid in understanding if the POCT HIV testing could be integrated with HCV testing

initiatives given the overlap of risks and social need, and services used.

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A limitation of the analysis could be conceptual overlap between the TDF and TFA frameworks.

There is a need to consider how these two frameworks can best be used concurrently, avoiding

redundancy. Similar issues have been raised in implementation studies, and it is suggested that

analysis plans outline the role of each framework used (44).

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These findings may reflect the specific context of one service and may not apply to other services

nationally, or internationally. Further research is recommended to generate evidence about the

generalisability of the results reported here.

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4.3. Conclusion

20 Uptake of HIV-POCT amongst service users was high. The location and approach to testing, along

with POCT itself was acceptable and feasible to service users and healthcare providers. The results

indicate that POCT targeted those considered to be at high risk of HIV infection and therefore could

provide benefits in terms of early detection of cases, better management of individuals, prevention

of onward transmission and early detection of a potential outbreak.

- 1 Further work is recommended to establish barriers and facilitators of POCTs across different
- 2 contexts and in different stages of the testing process. Healthcare providers require appropriate
- 3 training and supervision to offer and administer POCT in a sensitive manner.

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| TFA domain | TFA domain definition (24) | Example question from service user topic guide | Example question from service provider topic guide |
|-------------------------|---|---|--|
| Affective attitude | How an individual feels about the intervention | How did being offered the test make you feel? Prompt: Did you have any concerns? Prompt: Did the test provide reassurance? | Did you feel like it was your responsibility to offer the test? |
| Burden | The perceived amount of effort that is required to participate in the intervention | How convenient was the test? Prompt: How suitable a time and place was it for the test? | How do you feel about adding HIV point-of-care test to routine sessions? |
| Ethicality | The extent to which the intervention has good fit with an individuals value system | Did you think being offered the test by this team was appropriate? Prompt: Why do you think you were offered the test? | Do you think the test should also be offered to other people? |
| Intervention coherence | The extent to which the individual understands the intervention and how it works | What did you understand about the test and what it involved? Prompt: What information would you have liked to have? Prompt: What do you know about HIV? | What is your understanding of the HIV point-of-care test? |
| Opportunity costs | The extent to which benefits, profits or values must be given up to engage in the intervention | How much of a priority is your health in your day-to-day life? Prompt: What is more important for you? | Are there any other priorities that make the test difficult? |
| Perceived effectiveness | The extent to which the intervention is perceived as likely to achieve its purpose | How good a way of testing for HIV was the test? Prompt: Can you think of any other ways that may have worked better? | How effective do you think the test is? |
| Self-efficacy | The participant's confidence that they can perform the behaviour(s) required to participate in the intervention | Did you feel able to cope with the test and result? Prompt: Do you have any experience of similar tests? | How easy or difficult was it to administer the test? |
| TDF domain | TDF domain definition (25) | Service user example question | |
| Knowledge | An awareness of the existence of something | What did you understand about the test and what it involved? Prompt: What information would you have liked to have? Prompt: What do you know about HIV? | What other services are available? |

| Skills | An ability or proficiency acquired through practice | Did you feel able to cope with the test and result? Prompt: Do you have any experience of similar tests? | How did you communicate results? |
|--|---|---|--|
| Social/Professional role and identity | A coherent set of behaviours and displayed personal qualities of an individual in a social or work setting | Did you think being offered the test by HHT was appropriate? Prompt: Why do you think you were offered the test? | How much do you think this fits within your professional role? |
| Beliefs about capabilities | Acceptance of the truth, reality, or validity about an ability, talent, or facility that a person can put to constructive use | Did you feel able to cope with the test and result? Prompt: Do you have any experience of similar tests? | How easy or difficult was it to administer the test? |
| Optimism | The confidence that things will happen for the best or that desired goals will be attained | Did you think that finding out whether you had HIV was a good thing? Prompt: Why? | How effective do you think the test is? |
| Beliefs about consequences | Acceptance of the truth, reality, or validity about outcomes of a behaviour in a given situation | What did you think might happen because of having the test? Prompt: What do you think can be done if tests are positive? | Do you have any concerns about the test? |
| Reinforcement | Increasing the probability of a response by arranging a dependent relationship, or contingency, between the response and a given stimulus | What encouraged you to have the test? Prompt: How did you feel about having the choice to accept or decline the test? Prompt: How would you have felt about the test being given as standard? | Did you feel like it was your responsibility to offer the test? |
| Intention | A conscious decision to perform a behaviour or a resolve to act in a certain way | Would you consider having the test again in future? Prompt: What might affect your intention to have it? | How did you decide whether to offer the test? |
| Goals | Mental representations of outcomes or end states that an individual wants to achieve | How much of a priority is your health in your day-to-day life? Prompt: What is more important for you? | How much does the test make a difference to service users' health? |
| Memory, attention and decision process | The ability to retain information, focus selectively on aspects | Could you tell me about what led you to seek support from HHT today? | What are the advantages of offering the test? |

| | of the environment and choose between two or more alternatives | What influenced your decision to have the HIV point-of-care test? How much did it meet your expectations? | |
|-------------------------------------|---|--|--|
| Environmental context and resources | Any circumstance of a person's situation or environment that discourages or encourages the development of skills and abilities, independence, social competence, and adaptive behaviour | How convenient was the test? Prompt: How suitable a time and place was this service for the test? | Did the service have the resources it needed for the test |
| Social influences | Those interpersonal processes that can cause individuals to change their thoughts, feelings, or behaviours | Do you know other people who have had the test? Prompt: Would you discuss having had the test with other people? Prompt: Did how your service provider explained it influence your decision? | How did this fit with the team culture and ways of working? |
| Emotion | A complex reaction pattern, involving experiential, behavioural, and physiological elements, by which the individual attempts to deal with a personally significant matter or event | How did being offered the test make you feel? Prompt: Did you have any concerns? Prompt: Did the test provide reassurance? | How do you feel about adding HIV point-of-care test to routing sessions? |
| Behaviour regulation | Anything aimed at managing or changing objectively observed or measured actions | Was there anything you did on the day of the test that made you choose to have it? Prompt: How did attending affect your normal day-to-day routine? | Did you have to make any changes to your practice? |

| Age | Range 29-50, |
|-----------------------|--------------|
| | Median 40 |
| Gender | |
| Male | 64%, n=7 |
| Female | 36%, n=4 |
| Ethnicity | |
| White British | 91%, n=10 |
| White Other | 9%, n=1 |
| Housing status* | |
| Roofless: people | 36%, n=4 |
| living rough | |
| Insecure: people | 9%, n=1 |
| living in insecure | |
| accommodation | |
| Insecure: people | 36%, n=4 |
| receiving longer- | |
| term support (due | |
| to homelessness) | |
| Houseless: people in | 18%, n=2 |
| accommodation for | |
| the homeless | |
| HIV risk factors | |
| Bought or sold sex | 9%, n=1 |
| Sexual contact with | 73%, n=8 |
| someone known to | |
| be HIV positive (high | |
| risk partner) | |
| Injecting drug use | 91%, n=10 |
| Prison record | 54%, n=6 |
| Men who have sex | 0 |
| with men | |
| Reporting two or | 73%, n=8 |
| more HIV risk | |
| factors | |
| * Housing status | |
| defined by the | |
| European Typology | |
| of Homelessness | |
| and Housing | |
| Exclusion (FEANTSA, | |
| 2017) | |
| | |

| TDF | TFA | Theme | Facilitator/ Barrier | Quote from SU | Quote from SP |
|---|----------------------------|--|-------------------------|--|---|
| Knowledge | Intervention coherence | Understanding test purpose and process | Facilitator | "it's to see if you have HIV." Pt11 | "The [POCT] kits were easy to use" SP FG |
| Social/ professional role and Identity | Ethicality | Collective and individual responsibility | Facilitator | "I hope it can make them more willing to come and have one 'cause it could cure the people catching it." Pt 3 | "I think it does definitely fit with our role." FG |
| Beliefs about capabilities | Self-efficacy | Uncertainty over coping with positive results | Mixed | "if it did come back positive then the healthcare people are here to put the steps for you to sort it out quickly." Pt 9, | "I think as I say, we haven't had a reactive but we'll deal with that. We deal with far more essentially. Yes, we'll just deal with it as it comes along I think" FG. |
| | | Staff have the skills to administer test | Facilitator | "she knows I'm really hard to get blood out of. I may as well be of stone" Pt1 | "I guess that probably just became a bit more fluent yes" FG |
| Environmental and contextual resources | Burden | Ease and convenience of test | Facilitator | "It's all here in one place really so, you know, there's no reason why you can't access it." Pt5 | "it was an extra thing but actually if they were just coming in and almost like taking their blood pressure and you could explain it quicker than trying to write things down." FG |
| Optimism | Perceived effectiveness | Testing will have positive impact on the population | Facilitator | "'If I knew it was so easy to accept that I would have accepted it a long time ago." Pt 3 | "Probably a small chance where there is someone who we find a reactive test on. There is big potential for impact." FG. |
| Emotion | Affective attitude | Anxiety | Barrier | "A little bit scared, yeah. 'Cause I thought, 'What if I have got it?' I kept putting off getting the test done." Pt 5 | "I worried a little bit about was the whole idea that if it was reactive, we'd phone the sexual health clinic and somebody would see them and I was never convinced in my mind that's what would happen" FG |

| | | | " . | |
|--|--|-------------|--|--|
| | Stigma | Barrier | "Just me telling someone today before I was coming to do this interview thing it's like I had a bit of a look. You have to explain to them straightaway, 'Oh, there's nothing like that. It's just I had a test and things. I was offered it'" Pt8 | NA |
| Memory - attention and decision making processes | Choice and agency | Mixed | "I'd be on the back foot because it's my life at the end of the day. You know. I think that's down to me ultimately." Pt 6 | "not offer that to the patients on the day because actually we can't say. If we say 'Well I'm going to phone [SERVICE NAME] and get you an appointment today' because that's what we were told to do, I wouldn't do that now because we can't guarantee that they would get an appointment that day." FG |
| Beliefs about - consequences | Carrying out the test will influence the commissioning and staffing of the service | Facilitator | NA | "that would help them with their commissioning for staff and that kind of thing as well" FG |

