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Uptake and yield of HIV testing and counselling among children and adolescents in sub-Saharan Africa: a systematic review

Darshini Govindasamy, Rashida A Ferrand, Stephanie MS Wilmore, Nathan Ford, Saeed Ahmed, Hoviyeh Afnan-Holmes and Katharina Kranzer

Abstract

Introduction: In recent years children and adolescents have emerged as a priority for HIV prevention and care services. We conducted a systematic review to investigate the acceptability, yield and prevalence of HIV testing and counselling (HTC) strategies in children and adolescents (5 to 19 years) in sub-Saharan Africa.

Methods: An electronic search was conducted in MEDLINE, EMBASE, Global Health and conference abstract databases. Studies reporting on HTC acceptability, yield and prevalence and published between January 2004 and September 2014 were included. Pooled proportions for these three outcomes were estimated using a random effects model. A quality assessment was conducted on included studies.

Results and discussion: A total of 16,380 potential citations were identified, of which 21 studies (23 entries) were included. Most studies were conducted in Kenya (n = 5) and Uganda (n = 5) and judged to provide moderate (n = 15) to low quality (n = 7) evidence, with data not disaggregated by age. Seven studies reported on provider-initiated testing and counselling (PITC), with the remainder reporting on family-centred (n = 5), home-based (n = 5), outreach (n = 5) and school-linked HTC among primary schoolchildren (n = 1). PITC among inpatients had the highest acceptability (86.3%; 95% confidence interval [CI]: 65.5 to 100%), yield (12.2%; 95% CI: 6.1 to 18.3%) and prevalence (15.4%; 95% CI: 5.0 to 25.7%). Family-centred HTC had lower acceptance compared to home-based HTC (51.7%; 95% CI: 10.4 to 92.9% vs. 84.9%; 95% CI: 74.4 to 95.4%) yet higher prevalence (100%), yield (12.2%; 95% CI: 6.1 to 18.3%) and prevalence (15.4%; 95% CI: 5.0 to 25.7%). Family-centred HTC had lower acceptance compared to home-based HTC (51.7%; 95% CI: 10.4 to 92.9% vs. 84.9%; 95% CI: 74.4 to 95.4%) yet higher prevalence (8.4%; 95% CI: 3.4 to 13.5% vs. 3.0%; 95% CI: 1.0 to 4.9%). School-linked HTC showed poor acceptance and low prevalence.

Conclusions: While PITC may have high test acceptability priority should be given to evaluating strategies beyond healthcare settings (e.g. home-based HTC among families) to identify individuals earlier in their disease progression. Data on linkage to care and cost-effectiveness of HTC strategies are needed to strengthen policies.

Keywords: adolescents; children; HIV testing and counselling; sub-Saharan Africa.

To access the supplementary material to this article please see Supplementary Files under Article Tools online.

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prevalence of different HTC strategies in children and adolescents in SSA.

Methods
Search strategy and study inclusions
The search strategy aimed to identify evidence from randomized and non-randomized trials, prospective and retrospective cohort studies, cross-sectional studies and programme evaluations that reported on HTC among children and adolescents (5 to 19 years) with sufficient data to calculate prevalence of new HIV-positive diagnoses. Studies were omitted if participants did not receive their test results (i.e. anonymized surveys) or if the age range did not overlap with the targeted age range (5 to 19 years) by at least three years. Studies conducted in antenatal settings as part of the prevention of mother-to-child transmission (PMTCT) and in inpatient, outpatient, STI and TB clinics were only included for full text review if the abstract indicated that the age range overlapped with the targeted age range (5 to 19 years) by at least three years. The search was limited to studies conducted in SSA and published between 1 January 2004 and 30 September 2014, with no language restrictions. See protocol (Supplementary file 1) and PRISMA checklist attached (Supplementary file 2).

An electronic search was conducted on MEDLINE, EMBASE and Global Health using a compound search strategy (Supplementary file 3). A checklist of known studies was used to ensure that our search strategy captured all relevant studies. In addition, abstracts of all conferences of the International AIDS Society were screened from 2010 to 2013 to identify studies that may have been recently completed but not yet published. Reference lists of all eligible studies and systematic reviews were searched for additional articles, and authors of potentially eligible and included studies were contacted to provide age-stratified data. Two attempts were made to contact authors.

Data extraction
All references were imported into EndNote, and titles and abstracts were screened independently by two investigators (KK, DG). Full texts of potential studies were then obtained and the inclusion criteria applied. Final study inclusion was based on consensus between investigators (DG, KK). Data was then entered from each selected study onto a standardized data extraction form (DG) and cross-checked (KK). The following variables were extracted: study design, study setting, HTC strategy, type of HIV screening test used, number of participants who were offered HTC, number of participants who accepted HTC and number of participants testing HIV positive.

Definition of outcomes
The following definitions were applied: 1) testing acceptance rate, the proportion of individuals who underwent HTC and received their test results of those eligible for HTC; 2) yield of new HIV-positive diagnoses, the proportion of individuals who were newly diagnosed HIV positive of those who were eligible for HTC; and 3) prevalence of new HIV-positive diagnoses, the proportion of individuals who were newly diagnosed HIV positive of those who underwent HTC.

Quality assessment
The quality of evidence among included published studies was assessed using standardized criteria which examined misclassification, selection and reporting bias by evaluating the following factors: description of HTC procedures, inclusion and exclusion criteria, sampling strategy, reporting of HTC outcome data, discussion of limitations and sub-group analyses performed. Study quality was classified accordingly: high (score 8 to 10), moderate (score 5 to 7) or low (score 4 to 0).

Data analysis
For each included study, the numbers of individuals eligible, tested and tested positive were used to estimate proportions and corresponding 95% confidence intervals. Data were then pooled and stratified by testing strategy. On initial analysis, significant heterogeneity was found between studies. Therefore the pooled proportions of individuals accepting testing and testing positive (and 95% confidence interval, CI) were estimated with a random effects model, weighting for the inverse of the variance. Data analyses were conducted using Stata 12 (StataCorp, College Station, TX, USA).

Results and discussion
A total of 16,380 deduplicated potentially relevant citations were identified, including three systematic reviews on HTC [5–7] that provided 263 references to be screened. Ninety-four abstracts were identified for full-text review (Figure 1). A total of 21 studies were potentially eligible as there was overlap in the age range, but the data presented in the publication were not sufficiently stratified by age to enable data extraction for the age group of interest. In addition, adolescents aged 15 to 19 were mainly grouped with the 20- to 24-year-olds. Authors of all of these studies were contacted, resulting in an additional four studies (five entries for analysis) being included in the review [8–11].

Modes of HIV testing
The 21 studies (23 entries for analysis) included in the review (Table 1) reported data across eight countries: Kenya (n = 5) [12–16], Uganda (n = 5) [11,17–19], Zimbabwe (n = 4) [20–23], South Africa (n = 3) [8,24,25], Tanzania (n = 3) [9,10], Malawi (n = 1) [26], Sudan (n = 1) [27] and Zambia (n = 1) [28] (Table 1). Seven studies employed provider-initiated testing and counselling (PTC) for either inpatient (n = 3) or outpatient (n = 4) settings. Six studies were conducted in the context of sero-prevalence surveys, of which two provided HTC in the home environment and four used a mobile or outreach approach. A further four studies reported data from mass testing campaigns using outreach or home-based strategies. A family-centred approach was used in five studies, whereby patients known to HIV services were asked to identify other members in their households at risk of HIV infection. Finally, one study reported results from a school-linked testing campaign among primary schoolchildren aged 5 to 11 years [23]. Test uptake was comparable in girls and boys except for one study conducted in an outpatient setting in South Africa [25]. Fourteen studies reported HIV prevalence stratified by gender; of those, six [17,19–22,25] did not find any difference, and the remaining eight
(8–10,12,14,24,26) found a higher prevalence in girls compared to boys. The majority of studies (n = 22) used rapid point-of-care (POC) testing; testing conducted before 2005 was performed in the laboratory and results were either returned to participants at their homes or participants were asked to return to local primary healthcare clinics (PHC) for their test results. Most published studies on PITC, home-based HTC and outreach HTC were of moderate quality (n = 15), with seven studies judged to be of low quality because of a high degree of selection bias due to the sampling strategy used (i.e. consecutive sampling) and HTC being performed at set times during the day (Supplementary file 4).

Interpretations of findings
We investigated the uptake and yield of HTC among children and adolescents in SSA, the region where 90% of the world’s HIV-infected children live [4]. One of the key findings of this review is the lack of evidence for HTC approaches that are targeted towards children and adolescents. The HTC strategies employed predominantly replicate strategies developed for adults, with little consideration of the specific barriers associated with HTC and the needs of this age group [20]. HIV test acceptance, yield and prevalence differed according to setting and strategy. Generally yield and prevalence are influenced by 1) the overall HIV prevalence in the target group; 2) the refusal rate and 3) whether or not refusal is associated with HIV risk. Yield and prevalence are similar when refusal rates are low. The yield takes into account the refusal rate and thus is appropriate for comparing different testing strategies. The prevalence determines the number needed to test in order to diagnose a new case of HIV and has cost and resource implications.

The most common HTC strategy was healthcare facility-based testing, which in general reported a high uptake of HTC, particularly in hospital inpatient settings. The HIV yield and prevalence were also high, underscoring the importance of implementing routine HTC in healthcare facilities in high HIV prevalence settings. Reported barriers to PITC include lack of clear guidelines around consent procedures, prioritization of HTC within PMTCT programmes over testing of
<table>
<thead>
<tr>
<th>Author, Year</th>
<th>Country, region</th>
<th>Year of the study</th>
<th>Setting</th>
<th>Context</th>
<th>HTC strategy</th>
<th>Testing method</th>
<th>Eligible age group</th>
<th>Median or mean age</th>
<th>Proportion female</th>
<th>Number offered testing</th>
<th>Number accepted testing</th>
<th>Acceptance rate</th>
<th>Total testing</th>
<th>HIV positive</th>
<th>Yielda</th>
<th>Total Prevaleb</th>
<th>Quality of evidence</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ferrand et al. (2010)</td>
<td>Zimbabwe, Harare</td>
<td>2007 to 2008</td>
<td>Tertiary hospital, urban</td>
<td>Inpatient</td>
<td>PITC</td>
<td>POC testing</td>
<td>10 to 18 yrs</td>
<td>13 yrs</td>
<td>0.43</td>
<td>215</td>
<td>197</td>
<td>91.6%</td>
<td>50</td>
<td>23.3%</td>
<td>25.4%</td>
<td>No significant association between sex and HIV status ($p = 0.25$)</td>
<td>Moderate</td>
</tr>
<tr>
<td>Wanyenze et al. (2010)</td>
<td>Uganda, western to 2008</td>
<td>2005</td>
<td>Tertiary hospital, urban</td>
<td>Inpatient</td>
<td>PITC</td>
<td>POC testing</td>
<td>15 to 17 yrs</td>
<td>--</td>
<td>0.47</td>
<td>148</td>
<td>141</td>
<td>95.3%</td>
<td>--</td>
<td>10</td>
<td>6.8%</td>
<td>7.1%</td>
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<tr>
<td>Abbas et al. (2010)</td>
<td>Sudan, Khartoum</td>
<td>2007 to 2008</td>
<td>Tertiary hospital, urban</td>
<td>Inpatient</td>
<td>PITC</td>
<td>POC testing</td>
<td>15 to 17 yrs</td>
<td>5 yrs</td>
<td>0.42</td>
<td>127</td>
<td>106</td>
<td>83.5%</td>
<td>--</td>
<td>6</td>
<td>4.7%</td>
<td>5.7%</td>
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<tr>
<td>Kankasa et al. (2009)</td>
<td>Zambia, Lusaka</td>
<td>2006 to 2007</td>
<td>Tertiary hospital, urban</td>
<td>Inpatient</td>
<td>PITC</td>
<td>POC testing</td>
<td>6 to 18 yrs</td>
<td>--</td>
<td>--</td>
<td>1785</td>
<td>1060</td>
<td>59.4%</td>
<td>--</td>
<td>248</td>
<td>13.9%</td>
<td>23.4%</td>
<td>--</td>
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<td>Ramirez-Avila et al. (2012)</td>
<td>South Africa, Durban (KwaZulu-Natal)</td>
<td>2008 to 2009</td>
<td>Secondary hospital, urban</td>
<td>Outpatient</td>
<td>PITC</td>
<td>POC testing</td>
<td>12 to 17 yrs</td>
<td>--</td>
<td>0.55</td>
<td>956</td>
<td>389</td>
<td>40.7%</td>
<td>49.0%</td>
<td>30.0%</td>
<td>$p &lt; 0.01$</td>
<td>62</td>
<td>6.5%</td>
</tr>
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<td>Zimbabwe, Harare</td>
<td>2013</td>
<td>PHC</td>
<td>Outpatient</td>
<td>PITC</td>
<td>POC testing</td>
<td>6 to 15 yrs</td>
<td>9 yrs IQR: 7 to 11</td>
<td>0.47</td>
<td>2151</td>
<td>1534</td>
<td>71.3%</td>
<td>No significant association between sex and acceptance rate</td>
<td>82</td>
<td>3.8%</td>
<td>5.3%</td>
<td>No significant association between sex and HIV status</td>
</tr>
<tr>
<td>Ferrand et al. (2010)</td>
<td>Zimbabwe, Harare</td>
<td>2009</td>
<td>PHC and ANC clinics, peri-urban</td>
<td>Outpatient</td>
<td>PITC (n = 506), ANC (n = 88)</td>
<td>POC testing</td>
<td>10 to 18 yrs</td>
<td>APC: 14 yrs ANC: 17 yrs</td>
<td>APC: 0.58</td>
<td>594</td>
<td>573</td>
<td>96.5%</td>
<td>--</td>
<td>75</td>
<td>12.6%</td>
<td>13.1%</td>
<td>For APC, no significant association between sex and HIV status ($p = 0.35$)</td>
</tr>
<tr>
<td>Mongare et al. (2013)</td>
<td>Kenya</td>
<td>2009 to 2012</td>
<td>Primary healthcare facilities, peri-urban</td>
<td>Outpatient</td>
<td>Family-centred</td>
<td>POC testing</td>
<td>0 to 15 yrs</td>
<td>--</td>
<td>22,688</td>
<td>7382</td>
<td>32.5%</td>
<td>--</td>
<td>839</td>
<td>3.7%</td>
<td>11.4%</td>
<td>--</td>
<td>N/A</td>
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<td>Kenya</td>
<td>2007 to 2009</td>
<td>Primary healthcare facilities, urban</td>
<td>Outpatient</td>
<td>Family-centred</td>
<td>POC testing</td>
<td>0 to 15 yrs</td>
<td>--</td>
<td>--</td>
<td>484</td>
<td>276</td>
<td>57.0%</td>
<td>--</td>
<td>50</td>
<td>10.3%</td>
<td>18.1%</td>
<td>--</td>
</tr>
<tr>
<td>Were et al. (2006)</td>
<td>Uganda, districts of Busia</td>
<td>2003 to 2004</td>
<td>Community, urban, rural</td>
<td>Home-based</td>
<td>Family-centred</td>
<td>Laboratory testing, results provided at people’s homes</td>
<td>6 to 10 yrs</td>
<td>--</td>
<td>0.48</td>
<td>604</td>
<td>602</td>
<td>99.6%</td>
<td>99.0%</td>
<td>99.0%</td>
<td>23</td>
<td>3.8%</td>
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<td>Were et al. (2006)</td>
<td>Uganda, districts of Tororo and Busia</td>
<td>2003 to 2004</td>
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<td>Home-based</td>
<td>Family-centred</td>
<td>Laboratory testing, results provided at people’s homes</td>
<td>11 to 17 yrs</td>
<td>--</td>
<td>0.48</td>
<td>737</td>
<td>734</td>
<td>99.6%</td>
<td>99.0%</td>
<td>99.0%</td>
<td>12</td>
<td>1.6%</td>
<td>1.6%</td>
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<td>Country, region</td>
<td>Year of study</td>
<td>Setting</td>
<td>Context</td>
<td>HTC strategy</td>
<td>Testing method</td>
<td>Eligible age group</td>
<td>Median or mean age</td>
<td>Proportion female</td>
<td>Number offered testing</td>
<td>Number accepted testing</td>
<td>Acceptance rate</td>
<td>Total testing HIV positive</td>
<td>Yield</td>
<td>Prevalenceb</td>
<td>Quality of evidence</td>
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<td>Lugada et al. (2010)</td>
<td>Uganda, southeastern to 2007</td>
<td>Community, rural, urban</td>
<td>Home-based</td>
<td>Family-centred HTC</td>
<td>POC testing</td>
<td>6 to 14 yrs</td>
<td>–</td>
<td>0.53</td>
<td>1779</td>
<td>1055</td>
<td>59.3% 58.2% 60.6%</td>
<td>24</td>
<td>1.3% 2.3% 2.7% 1.8%</td>
<td>Moderate</td>
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<td>PHC Outpatient</td>
<td>Community, rural</td>
<td>Home-based HTC</td>
<td>POC testing</td>
<td>6 to 14 yrs</td>
<td>–</td>
<td>0.53</td>
<td>979</td>
<td>96</td>
<td>9.8% 8.6% 11.1%</td>
<td>9</td>
<td>0.9% 9.4% 8.9% 9.8%</td>
<td>Moderate</td>
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<td>Naik et al. (2012)</td>
<td>South Africa, Sisonke District to 2011</td>
<td>Community, rural</td>
<td>Testing campaign</td>
<td>Home-based HTC</td>
<td>POC testing</td>
<td>14 to 19 yrs</td>
<td>–</td>
<td>0.65</td>
<td>1011</td>
<td>867</td>
<td>85.8% 86.9% 85.6%</td>
<td>32</td>
<td>3.2% 3.7% 5.5% 0.3%</td>
<td>Moderate</td>
<td></td>
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<tr>
<td>Wachira et al. (2014)</td>
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<td>Community, urban/rural</td>
<td>Testing campaign</td>
<td>Home-based HTC</td>
<td>POC testing</td>
<td>13 to 18 yrs</td>
<td>15.3 yrs</td>
<td>0.50</td>
<td>34,607</td>
<td>34,410</td>
<td>99.4% No significant association between sex and acceptance rate</td>
<td>162</td>
<td>0.5% Females had greater odds of testing HIV positive</td>
<td>Low Moderate</td>
<td></td>
<td></td>
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<tr>
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<td>Kenya, western</td>
<td>Community, urban</td>
<td>Testing campaign</td>
<td>Home-based HTC</td>
<td>POC testing</td>
<td>18 mo to 13 yrs</td>
<td>–</td>
<td>0.48</td>
<td>2289</td>
<td>1294</td>
<td>56.5% 58.0% 55.2%</td>
<td>60</td>
<td>2.6% 4.6% – –</td>
<td>Moderate</td>
<td></td>
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<tr>
<td>Dalal et al. (2013)</td>
<td>Kenya, Lwak (Nyanza Province), Kibera (Nairobi)</td>
<td>Community, urban/rural</td>
<td>Sero-prevalence survey</td>
<td>Home-based HTC</td>
<td>POC testing</td>
<td>&lt; 13 yrs</td>
<td>–</td>
<td>0.49</td>
<td>1234</td>
<td>1190</td>
<td>96.4% 96.8% 96.0%</td>
<td>136</td>
<td>11.0% 11.4% 12.9% 10.0%</td>
<td>Moderate</td>
<td></td>
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<td>Angotti et al. (2009)</td>
<td>Malawi, North, Central and South Regions</td>
<td>Community, rural</td>
<td>Sero-prevalence survey</td>
<td>HTT</td>
<td>laboratory testing, results provided at people's homes</td>
<td>2004, POC testing 2006</td>
<td>15 to 19 yrs</td>
<td>–</td>
<td>0.49</td>
<td>1076</td>
<td>1007</td>
<td>93.6% 93.5% 93.7%</td>
<td>6</td>
<td>0.6% 0.6% 1.0% 0.2%</td>
<td>Moderate</td>
<td></td>
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<tr>
<td>Kranzer et al. (2011)</td>
<td>South Africa, Cape Town</td>
<td>Community, peri-urban</td>
<td>Sero-prevalence survey</td>
<td>Outreach (Mobile clinic with home-based invitation)</td>
<td>POC testing</td>
<td>15 to 19 yrs</td>
<td>–</td>
<td>–</td>
<td>140</td>
<td>119</td>
<td>85.0% – – – –</td>
<td>3</td>
<td>2.1% 2.5% – –</td>
<td>Females had greater odds of testing HIV positive</td>
<td>Moderate</td>
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<td>Tanzania, Mwanza Region (Lake zone)</td>
<td>Community, urban, peri-urban</td>
<td>Sero-prevalence survey</td>
<td>Outreach HTC at central site (opt in)</td>
<td>POC testing</td>
<td>15 to 19 yrs</td>
<td>–</td>
<td>0.55</td>
<td>1302</td>
<td>786</td>
<td>60.4% 61.4% 59.1%</td>
<td>7</td>
<td>0.5% 0.9% 1.4% 0.3%</td>
<td>Moderate</td>
<td></td>
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<tr>
<td>Author, Year</td>
<td>Country, region</td>
<td>Study period</td>
<td>Setting</td>
<td>Context</td>
<td>HTC strategy</td>
<td>Testing method</td>
<td>Eligible age group</td>
<td>Median or mean age</td>
<td>Proportion female</td>
<td>Number offered testing</td>
<td>Number accepted testing</td>
<td>Acceptance rate</td>
<td>Total testing HIV positive</td>
<td>Yield(^a)</td>
<td>Total Female</td>
<td>Male</td>
<td>Female</td>
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</tr>
<tr>
<td>Baisley et al. (2012)</td>
<td>Tanzania, Mwanza Region (Lake zone)</td>
<td>2007 to 2008</td>
<td>Community, urban, peri-urban</td>
<td>Sero-prevalence survey</td>
<td>Outreach HTC at central site (opt out)</td>
<td>POC testing</td>
<td>15 to 19 yrs</td>
<td>–</td>
<td>0.57</td>
<td>1223</td>
<td>1103</td>
<td>90.2%</td>
<td>91.1%</td>
<td>89.0%</td>
<td>11</td>
<td>0.9%</td>
<td>1.0%</td>
</tr>
<tr>
<td>Isingo et al. (2012)</td>
<td>Tanzania, Kiesa ward (Mwanza Region)</td>
<td>2003 to 2004</td>
<td>Community, rural</td>
<td>Sero-prevalence survey</td>
<td>Outreach HTC at a central site</td>
<td>Laboratory testing, return for results in 2003/4 and POC testing in 2006/7</td>
<td>15 to 19 yrs</td>
<td>–</td>
<td>0.50</td>
<td>2244</td>
<td>223</td>
<td>9.9%</td>
<td>10.8%</td>
<td>9.1%</td>
<td>17</td>
<td>0.8%</td>
<td>7.6%</td>
</tr>
<tr>
<td>Chamie et al. (2014)</td>
<td>Uganda</td>
<td>2012</td>
<td>Community, rural</td>
<td>Testing campaign</td>
<td>Outreach HTC</td>
<td>POC testing</td>
<td>10 to 19 yrs</td>
<td>–</td>
<td>–</td>
<td>1762</td>
<td>998</td>
<td>56.6%</td>
<td>–</td>
<td>–</td>
<td>5</td>
<td>0.30%</td>
<td>0.50%</td>
</tr>
<tr>
<td>Bandason et al. (2013)</td>
<td>Zimbabwe, Harare</td>
<td>2010</td>
<td>Peri-urban</td>
<td>Schools</td>
<td>School-linked HTC</td>
<td>POC testing</td>
<td>5 to 9 yrs</td>
<td>–</td>
<td>0.54</td>
<td>2273</td>
<td>22</td>
<td>1.0%</td>
<td>1.0%</td>
<td>0.9%</td>
<td>1</td>
<td>0.04%</td>
<td>4.5%</td>
</tr>
<tr>
<td></td>
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<td></td>
<td></td>
<td></td>
<td></td>
<td>10 to 11 yrs</td>
<td>–</td>
<td>0.56</td>
<td>1334</td>
<td>11</td>
<td>0.8%</td>
<td>0.8%</td>
<td>0.9%</td>
<td>1</td>
<td>0.1%</td>
<td>9.0%</td>
</tr>
</tbody>
</table>

\(^a\)HIV prevalence among all eligible; \(^b\)HIV prevalence among all tested. ANC, antenatal clinic; APC, acute primary care; HTC, HIV testing and counselling; IQR, interquartile ratio; PITC, provider-initiated testing and counselling; PHC, primary healthcare centre; VCT, voluntary counselling and testing; POC, point of care; N/A, not applicable.
children and adolescents and perceived lack of skills among healthcare providers to discuss HTC with children, adolescents and their guardians [20]. Despite the relatively high acceptance and yield of PITC as well as linkage to care (95%) [20], the crucial caveat is that this strategy mainly identifies children in inpatient settings when they are symptomatic and likely to have advanced disease [21,22]. In contrast, community-based HTC approaches have the potential to diagnose children at an earlier stage of infection, as they do not rely on individuals presenting with symptoms. However, many studies reporting on such approaches tend to exclude children and adolescents [6].

Only one study, conducted in Zimbabwe, evaluated school-linked HTC among primary schoolchildren, which had the lowest acceptance rate compared to the other HTC strategies (1%) [23]. Key barriers described in this study were parents’ concern about confidentiality, stigma, inadvertent disclosure of their own HIV diagnosis and its likely adverse consequences. Healthcare workers were reluctant to test children (the majority of whom were orphaned) who had no legally-defined guardians, a concern which has also emerged in facility-based PITC [20]. South Africa is planning an extensive high school-linked HTC campaign, which forms part of the “basket of services” offered by the new Integrated School Health Program [3]. A recent qualitative study found that parents were generally in favour of school-linked HTC. However, they were not aware of their parental limitations in terms of the South African Children’s Act, which acknowledges that consent for an HIV test may be given by the child, if the child is over 12 years of age [29]. The inability to consent to HTC due to legal age restrictions in other countries in the region poses a challenge to school-linked testing programmes. This situation poses an additional barrier to HTC in Africa, where minors often live in extended families with no clearly defined guardian, as parents may have died or be absent for work [30–32]. School-linked HTC warrants further rigorous investigation into the appropriate age group to target, age at which one can provide consent and methods of subsequent referral to HIV care, given the high HIV prevalence reported in a South African survey among high school learners (4.7%), particularly among teenage girls (7.7%) [33].

Four studies reported on family-centred testing whereby an adult patient on ART (the parent) acts as the index case and triggers testing of the whole household, including children and adolescents at risk [15–17]. This method of case finding has been employed in the context of tuberculosis contact tracing for decades [34,35]. The vast majority of HIV infections in SSA are acquired sexually and vertically, resulting in strong spatial clustering of HIV within households [36]. Family-centred HTC was either offered to invited individuals at PHC [15,16,19] or through home-based HTC [17,19], which might explain the difference in uptake. The Kenyan family-centred model reported a relatively low acceptance rate (32.5 to 57.0%), but a high prevalence among adolescents tested (11.4 to 18.1%). The former might be due to logistical problems such as transport, a challenge often reported in the context of tuberculosis contact tracing [37–39]. This could be addressed by testing family members

<table>
<thead>
<tr>
<th>Strategy</th>
<th>Acceptance rate (%)</th>
<th>Yield (%)</th>
<th>Prevalence (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>PITC, inpatient (n=4)</td>
<td>59.4 to 95.3</td>
<td>86.3 (65.5 to 100)</td>
<td>99.6 (5.0 to 25.2)</td>
</tr>
<tr>
<td>PITC, outpatient (n=3)</td>
<td>40.7 to 96.5</td>
<td>68.5 (41.1 to 97.9)</td>
<td>99.8 (5.3 to 15.9)</td>
</tr>
<tr>
<td>Family centred HTC (n=5)</td>
<td>96.8 to 99.6</td>
<td>78.5 (41.6 to 97.9)</td>
<td>99.8 (5.3 to 15.9)</td>
</tr>
<tr>
<td>Home-based HTC (n=5)</td>
<td>96.8 to 99.6</td>
<td>78.5 (41.6 to 97.9)</td>
<td>99.8 (5.3 to 15.9)</td>
</tr>
<tr>
<td>Outreach (n=5)</td>
<td>96.8 to 99.6</td>
<td>78.5 (41.6 to 97.9)</td>
<td>99.8 (5.3 to 15.9)</td>
</tr>
</tbody>
</table>

CI, confidence interval; HTC, HIV testing and counselling; PITC, provider-initiated testing and counselling.

Table 2. Ranges and summary estimates

<table>
<thead>
<tr>
<th>Strategy</th>
<th>Acceptance rate (%)</th>
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<th>Prevalence (%)</th>
</tr>
</thead>
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</tr>
<tr>
<td>Family centred</td>
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</tr>
<tr>
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</tr>
</tbody>
</table>

CI, confidence interval; HTC, HIV testing and counselling; PITC, provider-initiated testing and counselling.
in the household rather than making them come to the health-care facility. Home-based-family-centred HTC resulted in a six times increase in test uptake among 6- to 14-year-olds in Uganda compared to family-centred HTC provided at PHCs [19].

Outreach HTC strategies had a low acceptance, yield and prevalence across all HTC strategies. The wide range of acceptance rates among studies investigating outreach HTC might be explained by differences in testing methods (POC vs. laboratory based testing) and differences in denominators. The study by Chamie et al. [11] was conducted as part of a testing campaign and denominators were estimated from a previous household census. Thus individuals not at home during the testing campaign were counted as eligible, resulting in an underestimation of test acceptance. Mobile HTC and outreach strategies have been successfully implemented in many SSA settings [40–42]. However, only one study included in this review used an outreach approach in the context of a testing campaign [11]. All other studies using outreach or mobile services were conducted as part of community prevalence surveys and as such were not representative. However, a recent cluster randomized trial investigating the effect of mobile community-based HTC resulted in a significant increase in testing rates among 16- to 17-year-olds [43].

Strength and limitations
This review has several strengths and limitations. We used an extensive search strategy including multiple databases and conference abstracts without language restrictions. Anticipating that data on adolescents might be reported as part of paediatric and adult studies, we included adult and paediatric studies in our search strategy and contacted authors of potentially eligible studies to obtain data disaggregated by age. However, additional data was only obtained for one-fifth of potentially eligible studies. As with any systematic review, this review is subject to publication bias. Specifically testing strategies with low acceptability might be less likely to be published in peer-reviewed literature. Due to paucity of data, this review was unable to assess the differences in outcomes disaggregated by age and whether HIV was acquired vertically (i.e. perinatally infected long-term survivors) or horizontally. Six of fourteen studies reported a similar HIV prevalence in girls compared to boys, which might indicate that those studies mainly targeted vertically infected children. Finally, none of the included studies assessed linkage to care, cost or cost-effectiveness, which are important factors policy makers need to consider when deciding which strategies to implement.

Conclusions
Achieving universal coverage of HTC for key populations in SSA under scarce resource constraints will require the implementation of innovative, effective and economically efficient population-based HTC strategies which can be readily brought to scale [4]. Data from our review indicate that HTC approaches delivered within communities outside of a healthcare facility (i.e. home-based, family-centred and outreach) have a high acceptance among this priority age group. Additionally these strategies have the potential to identify individuals early in their stage of HIV infection. However, there is a paucity of data on HTC strategies that extend beyond the healthcare facility for children and adolescents, particularly in areas where data are scarce such as school-linked, family-centred and mobile HTC. Moreover it is necessary to assess linkage to care and the cost-effectiveness of these different HTC approaches. Thus further evaluations are required prior to policy makers and programme managers planning for their scale-up. Furthermore, qualitative studies establishing the barriers to testing for this age group should be encouraged. Those barriers are likely to be specific to both the testing strategy and the setting [20,44].

Authors’ affiliations
1 Health Systems Research Unit, South African Medical Research Council, Cape Town, South Africa; 2 Department of Clinical Research, London School of Hygiene and Tropical Medicine, London, United Kingdom; 3 Biomedical Research and Training Institute, Harare, Zimbabwe; 4 Centre for Clinical Microbiology, University College London, London, United Kingdom; 5 HIV/AIDS Department, World Health Organization, Geneva, Switzerland; 6 Tingathe Outreach Program, Baylor College of Medicine Children’s Foundation Malawi, Lilongwe, Malawi; 7 Department of Medicine, Baylor College of Medicine, Houston, TX, USA

Competing interests
The authors have declared that no competing interests exist.

Authors’ contributions
KK conceptualized the review, conducted the search and drafted the proposal. DG and KK conducted the screening, data extraction and analysis. DG, KK and RAF prepared the first draft of the manuscript. DG, KK, RAF, SMSW, NF, SA and HA-H reviewed and commented on all subsequent drafts of the manuscript including the final draft. All authors approved the final version of this manuscript.

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Ethics
Ethical approval was not required for this work.

References


