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Effects of community-based antiretroviral therapy initiation models on HIV treatment outcomes: A systematic review and metaanalysis

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Abstract

Background

Antiretroviral therapy (ART) initiation in the community and outside of a traditional health facility has the potential to improve linkage to ART, decongest health facilities, and minimize structural barriers to attending HIV services among people living with HIV (PLWH). We conducted a systematic review and meta-analysis to determine the effect of offering ART initiation in the community on HIV treatment outcomes.

Methods and findings

We searched databases between 1 January 2013 and 22 February 2021 to identify randomized controlled trials (RCTs) and observational studies that compared offering ART initiation in a community setting to offering ART initiation in a traditional health facility or alternative community setting. We assessed risk of bias, reporting of implementation outcomes, and real-world relevance and used Mantel–Haenszel methods to generate pooled risk ratios (RRs) and risk differences (RDs) with 95% confidence intervals. We evaluated heterogeneity qualitatively and quantitatively and used GRADE to evaluate overall evidence certainty. Searches yielded 4,035 records, resulting in 8 included studies—4 RCTs and 4 observational studies—conducted in Lesotho, South Africa, Nigeria, Uganda, Malawi, Tanzania, and Haiti—a total of 11,196 PLWH. Five studies were conducted in general HIV populations, 2 in key populations, and 1 in adolescents. Community ART initiation strategies included community-based HIV testing coupled with ART initiation at home or at community venues; 5 studies maintained ART refills in the community, and 4 provided refills at the health facility. All studies were pragmatic, but in most cases provided additional resources. Few studies reported on implementation outcomes. All studies showed higher ART uptake in community 300342–104). The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

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Abbreviations: ART, antiretroviral therapy; FSW, female sex worker; PLWH, people living with HIV; PWID, people who inject drugs; RD, risk difference; RR, risk ratio; RCT, randomized controlled trial; WHO, World Health Organization. initiation arms compared to facility initiation and refill arms (standard of care) (RR 1.73, 95% CI 1.22 to 2.45; RD 30%, 95% CI 10% to 50%; 5 studies). Retention (RR 1.43, 95% CI 1.32 to 1.54; RD 19%, 95% CI 11% to 28%; 4 studies) and viral suppression (RR 1.31, 95% CI 1.15 to 1.49; RD 15%, 95% CI 10% to 21%; 3 studies) at 12 months were also higher in the community-based ART initiation arms. Improved uptake, retention, and viral suppression with community ART initiation were seen across population subgroups—including men, adolescents, and key populations. One study reported no difference in retention and viral suppression at 2 years. There were limited data on adherence and mortality. Social harms and adverse events appeared to be minimal and similar between community ART initiation and standard of care. One study compared ART refill strategies following community ART initiation (community versus facility refills) and found no difference in viral suppression (RD -7%, 95% CI -19% to 6%) or retention at 12 months (RD -12%, 95% CI -23% to 0.3%). This systematic review was limited by few studies for inclusion, poor-quality observational data, and short-term outcomes.

Conclusions

Based on data from a limited set of studies, community ART initiation appears to result in higher ART uptake, retention, and viral suppression at 1 year compared to facility-based ART initiation. Implementation on a wider scale necessitates broader exploration of costs, logistics, and acceptability by providers and PLWH to ensure that these effects are reproducible when delivered at scale, in different contexts, and over time.

Author summary

Why was this study done?

- Over the last decade HIV services have increasingly moved out of the health facility and into the community through the provision of decentralized and differentiated HIV care.
- It remains unclear however whether initiating ART in a community setting will result in treatment and safety outcomes that are comparable to facility-based ART initiation.

What did the researchers do and find?

- We conducted a systematic review to identify studies where ART was initiated at community locations, including homes, mobile vans, or other community venues.
- We identified 8 studies (including 11,196 HIV-positive people), 7 of which were conducted in sub-Saharan Africa; 4 were randomized controlled trials and 4 were cohort studies.
- The methodological quality of the randomized controlled trials was high, but cohort data were of poorer quality. Studies were generally pragmatic in design, but implementation outcomes were infrequently reported.
- Based on meta-analysis of this limited dataset, it appeared that ART initiation in the community resulted in higher ART uptake, higher retention, and greater viral

suppression at 1 year compared to facility-based ART initiation and refill, among HIVpositive people offered ART. These findings were consistent across population subgroups and various implementation strategies.

• There were insufficient data on serious adverse events or mortality to draw firm conclusions on these outcomes.

What do these findings mean?

- Community ART initiation may result in better outcomes than ART initiation in traditional health facilities.
- To increase the robustness of these findings, high-quality implementation research conducted in diverse settings, exploring optimum combinations of community ART initiation and ART refill strategies over longer time periods, will be critical.

Introduction

Initiating antiretroviral therapy (ART) in the community setting and outside of traditional health facilities represents an innovative addition to "differentiated service delivery" models, which seek to offer a greater range of options that meet a diversity of patient needs in the global HIV service delivery enterprise. Community-based ART initiation has several potential bene-fits including reducing psychological and structural barriers that newly diagnosed HIV-positive people face in order to access a facility, as well as further decongesting crowded facilities themselves. With a shift over the last decade to rapidly initiating ART in those who test HIV-positive, and decentralizing and differentiating follow-up after treatment initiation in the facility, community ART initiation represents the next step toward more patient-centered services and may bridge the critical gap between testing and linkage to ART—a point in the HIV care cascade when many disengage from care [1–3].

Despite this rationale, few studies have explored the effect of community ART initiation on either short- or long-term outcomes. Community-based HIV testing, including mobile testing, self-testing, testing campaigns, workplace testing, and index testing, frequently shows higher coverage and uptake than traditional facility-based testing and has the ability to reach those underserved by routine facility-based testing, particularly men and key populations [4,5]. Community-based medication refill for patients stable on ART, such as distribution directly to patients' homes or to community pick-up locations and pharmacies, has generally demonstrated success [6,7]. ART initiation, however, has traditionally been reserved for facilities because of the perceived intensity of the encounter, but this assumption is not empirically supported. Qualitative data suggest that the act of going to a clinic is intimidating and confusing, particularly in environments where stigma is present—making community ART initiation a potentially important innovation [8].

A number of studies examining the effects of community ART initiation have been conducted, but synthesis and review are needed to appraise the quality of the data as well as assess the top-line evidence of the effect of this approach on immediate and medium-term outcomes. In addition, systematic reviews are required as a part of the guideline development processes led by the HIV department at the World Health Organization (WHO), and therefore an important step in the translation of evidence to practice. To explore the effect of initiating ART in a community setting on HIV treatment outcomes, we conducted a systematic review that additionally characterized features of community ART initiation strategies to inform policy and implementation.

Methods

Search strategy and selection criteria

The protocol for this systematic review was registered on PROSPERO (CRD42019130272) and followed PRISMA guidelines [9,10] (S1 Table). We searched the Cochrane Central Register of Controlled Trials (CENTRAL, published in the Cochrane Library), MEDLINE (PubMed), Embase (OVID), Africa-Wide Information and CINAHL (EBSCOhost), LILACS, and Web of Science Core Collection from 1 January 2013 until 15 April 2019; this start date was based on the 2013 WHO recommendation for decentralization of HIV treatment and care [11], and a previous systematic review that searched for studies published from 1996 to 2013 and found no published studies on community-based ART initiation prior to 2013 [12] (S1 Appendix). Searches were updated on 1 April 2020 and again on 22 February 2021.

We included randomized and non-randomized study designs that enrolled HIV-positive people of any age, conducted in low- and middle-income countries, and compared community ART initiation to facility-based ART initiation or to another community-based ART initiation strategy. We defined community ART initiation as initiation of ART outside of a traditional health facility or workplace health center, by any cadre of health staff. Community settings included, for example, mobile health services, community centers, and patients' homes. ART could be offered and initiated in the community and subsequently maintained in the community or at a health center. Our comparison arm was ART initiation in a traditional health facility; ART maintenance after initiation could occur within or outside of the health facility. No language or age restrictions were applied to the search. We additionally searched HIV/AIDS conferences including International AIDS Society (IAS) conferences and the Conference on Retroviruses and Opportunistic Infections (CROI) until 10 March 2021, as well as the reference lists of included studies and relevant systematic reviews. We also searched ClinicalTrials. gov and the WHO International Clinical Trials Registry Platform for ongoing studies.

Data extraction and methodological quality assessment

Abstract and full-text screening was done in duplicate, with discrepancies resolved by a third author. Data from included studies were abstracted by a single author (SAA) and verified by a second author (AAA). Data were extracted in a pre-piloted data extraction tool developed in Airtable (https://airtable.com)—a commercially available web-based relational database tool. We extracted key characteristics of each study, including (1) study location; (2) methods: study design, dates and duration of study and follow-up, and number and type of sites; (3) study population: number, age, sex, and inclusion/exclusion criteria; (4) intervention and comparator details; (5) outcomes: ART uptake, retention in care, viral suppression, mortality, adherence, and adverse events, extracted when possible with numerators, denominators, and/or measures of association; and (6) indicators of risk of bias. Any discrepancies were resolved by discussion among the authors. The Cochrane or Newcastle–Ottawa Scale tools were used to assess risk of bias [13,14].

Assessment of study implementation characteristics

We additionally characterized studies according to PRECIS-2 criteria for how pragmatic or explanatory included studies were—exploring eligibility assessments, recruitment procedures,

settings, organizational characteristics, flexibility in intervention delivery and adherence, follow-up, primary outcome reporting, and primary analyses [15]. Although the tool is optimized for randomized control trial (RCT) design, we applied PRECIS-2 concepts across all studies. We also explored reporting of implementation outcomes in primary and additional study publications across 8 domains: acceptability, adoption, appropriateness, implementation cost, feasibility, fidelity, adaptation, penetration, and sustainability [16].

Data synthesis and statistical analysis

For pairwise meta-analyses, we used random effects generic inverse variance meta-analytic models; we evaluated ART uptake (initiation) among all HIV-positive individuals, and for retention in care and viral suppression we assessed these outcomes among all who initiated ART. We determined risk ratios (RRs) and 95% confidence intervals for all outcomes and additionally present risk differences (RDs) where absolute effects were deemed valuable for interpretation. For cluster randomized trials we calculated the design effect using methods outlined in the Cochrane handbook (where dichotomous counts from each study are divided by the quantity $1 + [M - 1] \times ICC$, where M is the average cluster size and ICC is the intra-cluster correlation coefficient) to adjust estimates if we could not incorporate adjusted estimates directly from study publications. We used Mantel-Haenszel methods to generate pooled estimates of binary data. We evaluated heterogeneity qualitatively through examining forest plots and quantitatively through examination of the I^2 statistic. Between-study variance was evaluated using the Paule–Mandel estimator for Tau2 and the associated I^2 statistic [17]. We used subgroup analyses to explore heterogeneity. For each outcome, we generated forest plots overall and subgrouped where relevant by study design, population type, and implementation features. We conducted tests for subgroup differences to determine if cohort and RCT data could be pooled. R statistical software was used for all analyses [18].

Evidence appraisal

We evaluated the certainty/quality of the body of evidence contributing to the pooled effect estimate for each outcome using criteria recommended by the GRADE Working Group, including risk of bias, inconsistency, indirectness, imprecision, and other domains [19–22].

Results

Searches were conducted between 1 January 2013 and 22 February 2021 and yielded 4,035 abstracts for screening after deduplication. We identified 120 records for full-text screening and 3 possibly eligible ongoing studies; 105 studies were excluded with reasons, and 15 publications representing 8 studies were included in the qualitative synthesis, with 7 studies included in meta-analyses (Fig 1).

Four studies were RCTs, 2 individually randomized [23–25] and 2 cluster randomized [26,27]; 3 were cohort studies with a comparison arm [28–30], and 1 study was a single-arm cohort study [31]. Two studies were conducted in Nigeria [29,31] and Lesotho [23,27] each, and 1 each was conducted in South Africa [25], Uganda [32], Malawi [26], Tanzania [28], and Haiti [30]. Five studies were conducted in general populations, 2 included key population groups, and 1 included adolescents (Table 1).

Studies incorporated several differentiated service delivery features including task shifting/ sharing and changes in the location and frequency of services (<u>Table 2</u>). Community ART initiation was in most cases delivered by a small team including a nurse and counselor or a village health worker, but in some studies involved a larger team, including lab technicians, pharmacists, doctors, and additional community lay workers [28,29,31]. Within the community





Study	tudy Study Country Design		Setting	Number of HIV	Population type	Eligibility criteria for inclusion
Barnabas 2020 [24,25]	RCT	South Africa and Uganda	Rural regions	1,315	General HIV endemic communities	ART naïve, WHO stage 1–3, CD4 cell count > 100/µL, not pregnant or breastfeeding, negative TB symptom screen, normal renal function
Ibiloye 2018 [31]	Cohort (single arm)	Nigeria	Three districts in a central state	935	Several key populations: FSWs, MSM, PWID	Any CD4 cell count
Labhardt 2018 [23,33,35]	RCT	Lesotho	Rural northern region	274	General HIV endemic communities	ART naïve, WHO stage 1–3, not pregnant or breastfeeding, no chronic illness, CRAG negative
Amstutz 2021 [27]	RCT (cluster)	Lesotho	Rural northern region	257	General HIV endemic community	ART naïve; weight > 35 kg; no other chronic condition; physical, mental, and emotional ability to participate; remaining in district for HIV care
MacPherson 2014 [26]	RCT (cluster)	Malawi	Blantyre (urban center)	768	General HIV endemic communities	CD4 cell count < 350/µL or WHO stage 3 or 4 or pregnant or breastfeeding
Oladele 2018 [29]*	Cohort	Nigeria	Fourteen donor-funded high-HIV-burden districts (urban and rural)	6,270	General HIV endemic communities	CD4 cell count $<$ 500/µL or WHO stage 3 or 4
Reif 2017 [<u>30</u>]**	Cohort	Haiti	Port Au Prince (urban center)	760	Adolescents and young adults	Community care group: any CD4 count; historical cohort: CD4 cell count < 350/µL
Tun 2019 [28,36]	Cohort	Tanzania	High HIV prevalence (major trucking routes)	617	FSWs	Any CD4 count

Table 1. General description of included studies.

CRAG, cryptococcal antigen screening; FSW, female sex worker; MSM, men who have sex with men; PWID, people who inject drugs; RCT, randomized controlled trial; TB, tuberculosis.

*Data restricted to patients testing for HIV in the community in the post-intervention period for intervention and control areas.

**Unpublished data from conference abstract only.

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setting, 2 studies initiated ART in the home, and the remaining studies initiated ART at community venues (such as mobile vans, individual homes, and other community venues). All studies paired community ART initiation with community-based HIV testing strategies. Two studies included HIV self-testing strategies; in one this was the primary method of HIV testing [26], and in another HIV self-tests were distributed to those who declined testing and those away from home in a subset of participants [27]. The location of ART refill collection (after ART initiation) varied, with half of the studies having participants collect ART refills in the community [28,30,31] and the other half having participants collect ART refills at the health facility [23,26,29]; Once stabilized on ART, those receiving facility-based ART refills received ART refills every 3 months. All studies initiated ART rapidly, either on the same day as testing (5 studies) or within 7 days (3 studies). Six comparative studies compared community ART initiation to facility-based ART initiation, with a similar subsequent frequency of ART refills in all except 1 study [23], where refill frequency was monthly in the facility arm compared to every 3 months in the community arm. One comparative study compared different ART refill strategies across 2 community ART arms [27]. In several studies, community ART initiation strategies were also combined with additional demand creation [26,31], enhanced support strategies [28-30], or SMS reminders [27]-beyond what was offered in the facility-based initiation (standard of care) arm.

Table 2. Intervention strategy details.

Study	Community AR	Гgroup							Comparator
	Community ART initiation team	HIV testing site	ART initiation site	Time to ART start	Immediate follow-up	ART refill frequency	ART refill location	Additional support/ interventions beyond routine care	groups
Barnabas 2020 [24,25]	Nurse, lay provider, ± driver	Community	Mobile van	≤7 days	Phone call at 7 days, in person at 1 month	Every 3 months	Mobile van	Food parcels provided at each study visit, quarterly phone calls	Two comparisons: (1) facility ART initiation with facility ART maintenance, (2) facility ART initiation and community ART maintenance
Ibiloye 2018 [31]	Community facilitator, ART clinician, nurse, counselor, pharmacist, lab technician	Community	Choice of outreach venues (CBO offices, hotels/ guest houses)	Same day	Not described	Not described	Outreach venues in community	STI care	No comparator
Labhardt 2018 [23,33,35]	Nurse, counselor	Home	Home	Same day	12–16 days and 6 weeks at health facility	Every 3 months	Health facility	Medical care at health facility	Facility ART initiation and maintenance, monthly clinic visits
Amstutz 2021 [27]	Nurse, counselor	Home	Home	Same day	12–16 days at VHW's home	Every 3 months	VHW's home	Monthly ART reminder via SMS, viral load result triggered SMS	Home-based ART initiation and facility ART follow-up and maintenance
MacPherson 2014 [26]	Nurse, counselor	Home (HIV self-test)	Home	≤7 days	2–4 weeks at health facility	Every 3 months	Health facility	Demand creation (HIV self-test and home ART awareness campaigns)	Facility ART initiation and maintenance
Oladele 2018 [29]	Doctor, counselor, pharmacist, lab technician, nurse, community lay workers	Community	Point of identification in community	Same day	Phone calls/ SMS/home visit every 3 days for 2 weeks, first facility refill at 1 month	Every 3 months	Health facility	Community mobilization campaigns, task sharing between providers and lay counselors	Facility ART initiation and maintenance
Reif 2017 [30,37]	Nurse, peer educator	Facility and community	Community center	Same day	At 1 month	Monthly	Community center	Integrated clinical care, FP, STI care, peer support	Facility ART initiation and maintenance, monthly clinic visits, routine facility-based support groups
Tun 2019 [28,36]	Clinician, nurse, lab technician, peer educator	Community	Mobile tent, home	≤7 days	Not described	Not described	Mobile tent, home	STI care, condom distribution, FP, IPV care, TB screening, CaCx screening, escorted referrals	Facility ART initiation and maintenance

CBO, community-based organization; CaCx, cervical cancer; FP, family planning; IPV, intimate partner violence; STI, sexually transmitted infection; TB, tuberculosis; VHW, village health worker.

Macpherson 2014: 36% of those offered home ART in the home group chose facility ART initiation (64% selected home art initiation); Amstutz 2021: 6- and 12-month ART refill visits were at health facility due to viral load measurement.

Study	ART uptake/initiation among HIV+ individuals	Retention in care among ART initiators	Viral suppression among ART initiators	Adherence among ART initiators	Mortality among HIV + individuals
Barnabas 2020	NA	NA	Low risk	NA	NA
Labhardt 2018	Low risk	Low risk	Low risk	NA	Some concerns
Amstutz 2021 ¹	NA	Some concerns	Some concerns	NA	Some concerns
MacPherson 2014	Low risk	Low risk	NA	High risk	Some concerns
Oladele 2018 ²	Poor quality	NA	NA	NA	NA
Reif 2017 ³	Poor quality	Poor quality	NA	NA	NA
Tun 2019 ⁴	Poor quality	Poor quality	Poor quality	Poor quality	Poor quality

Table 3. Risk of bias assessments.

NA, not applicable.

Assessments based on Cochrane RoB 1 tool for randomized controlled trials (high risk, some concerns, low risk) or Newcastle–Ottawa Scale for cohort studies (poor quality, good quality). Detailed assessments available in <u>S2 Appendix</u>.

¹Allocation concealment was not possible; recruiting teams were aware of household assignments prior to recruitment; participants were, however, unaware of assignment during recruitment.

²Comparison group (facility referral for ART initiation) very small compared to home ART group; fundamental differences between community and facility art initiation groups (facility groups urban and with lower HIV prevalence); unadjusted numbers used in this analysis.

 3 Overall risk of bias influenced by different sources of comparison (historical cohort with CD4 cell count < $350/\mu$ L) and intervention group; unadjusted estimates used in analysis without controlling for any baseline characteristics; in addition, the comparative analysis remains unpublished.

⁴Comparison group drawn from a different region; ascertainment of exposure not described; unadjusted estimates used in analysis; baseline imbalance in the group characteristics; intervention group had substantially more newly diagnosed participants, which could affect uptake and retention.

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Data from RCTs were generally judged as having high methodological quality (low risk of bias), and observational data as having poor quality, as assessed by risk of bias tools (Table 3; S2 Appendix). Observational studies had several methodological limitations, primarily related to lack of comparability of study arms and inclusion of data that were not adjusted for baseline imbalances in the pairwise meta-analysis.

We used PRECIS-2 criteria to assess how pragmatic or explanatory included studies were (Tables 4 and S2): Overall, studies were highly pragmatic—conducted in real-world settings, with flexible approaches to intervention delivery and few additional measures to ensure adherence to ART beyond what would occur in routine practice. RCTs were on average less pragmatic than cohort studies. Study procedures in trials that appeared less applicable in the "real world" setting included the following: (1) more restrictive inclusion criteria that included, for example, CD4 count measurement, or exclusion of those who were pregnant, breastfeeding, or had chronic conditions [25,27,34]; (2) expertise and resources used to deliver the intervention, with large, well-trained, multidisciplinary teams initiating ART in the community in some studies [23,26], or the provision of food parcels at visits [25]; and (3) extensive patient follow-up, where tracing efforts appeared more rigorous than what may occur in routine practice [23] —approaches that may not be entirely reproducible at scale.

ART initiation

Seven studies reported on ART initiation after offer of community ART; ART uptake among those testing HIV-positive was high overall (85%), but there was substantial heterogeneity of measurement time points, and uptake across studies ranged from 37% to 100% (Fig 2). The lowest ART uptake (37%) was seen in a study where HIV self-testing was conducted and paired with community ART initiation [26]. A single-arm study conducted in key populations had an overall uptake of 77%, but this varied across key population subgroups, with uptake of

Study	Eligibility	Recruitment	Setting	Organization	Flexibility: Delivery	Flexibility: Adherence	Follow-up	Primary outcome	Primary analysis
	Who is selected to participate in the trial?	How are participants recruited into the trial?	Where is the trial being done?	What expertise and resources are needed to deliver the intervention?	How should the intervention be delivered?	What measures are in place to make sure participants adhere to the intervention?	How closely are participants followed up?	How relevant is it to participants?	To what extent are all data included?
Barnabas 2020	4	4	5	3	4	5	5	5	5
Ibiloye 2018	5	4	5	3	5	5	3	5	5
Labhardt 2018	4	3	5	3	4	4	2	5	5
Amstutz 2021	5	4	5	3	4	5	4	5	5
MacPherson 2014	5	4	4	3	3	5	4	5	5
Oladele 2018	5	5	3	3	5	5	5	5	4
Reif 2017	5	5	4	4	5	5	4	5	4
Tun 2019	5	3	5	4	4	5	3	5	5

Table 4. PRECIS-2 criteria/score.

Value of 5 (dark green) represents a very pragmatic approach and a value of 1 (yellow) represents a very explanatory approach. Detailed assessments presented in S2 Table.

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75% in female sex workers (FSWs), followed by men who have sex with men (68%) and people who inject drugs (PWID) (53%); partners of key populations in this study had generally high community ART uptake (93%) [31].

Five studies compared community ART initiation to facility-based ART initiation and refill (standard of care). The meta-analysis of these studies showed higher ART initiation when

Study	Population	Design	Initiated ART	HIV positive			Propo	ortion				95%CI	Weight
$\begin{array}{l} \textbf{3 months} \\ \text{Reif 2017} \\ \text{Labhardt 2018} \\ \text{Amstutz 2021} \\ \textbf{Overall effect} \\ \text{Heterogeneity: } J^2 = 81\% \left[42\%; 9 \right] \end{array}$	Adolescents General General 4%], τ^2 = 1.1147, p < 0.01	Cohort RCT RCT	50 134 228	50 137 257 444							1.00 0.98 0.89 0.96	[0.86; 1.00] [0.93; 0.99] [0.84; 0.92] [0.85; 0.99]	8.6% 13.7% 15.4% 37.7%
6 months MacPherson 2014 Ibiloye 2018 Overall effect Heterogeneity: J ² = 100% [99%;	General Key population 100%], $\tau^2 = 1.5551$, $\rho < 0.0$	RCT Cohort	181 724	490 935 1425		_	Ð				0.37 0.77 0.59	[0.33; 0.41] [0.75; 0.80] [0.20; 0.89]	15.6% 15.6% 31.2%
12 months Tun 2019 Oladele 2018 Overall effect Heterogeneity: J ² = 96% [89%; 9	Key population General 9%], $\tau^2 = 0.2849$, $p < 0.01$	Cohort Cohort	256 4000	309 5789 6098				*			0.83 0.69 0.76	[0.78; 0.87] [0.68; 0.70] [0.60; 0.87]	15.5% 15.6% 31.1%
Overall effect Heterogeneity: <i>I</i> ² = 98% [97%; 9 Test for subgroup differences: <i>p</i>	9%], $\tau^2 = 2.4431$, $p < 0.01$ = 0.03			7967	0	0.2	0.4	0.6	0.8	-	0.85	[0.62; 0.95]	100.0%

Fig 2. ART uptake in community ART initiation study arms, among HIV-positive and by outcome measurement time point. Tun 2019 was conducted in female sex workers; Ibiloye 2018 included female sex workers, men who have sex with men, people who inject drugs, and partners of individuals in these key populations.

Study	COM(e)	COM(n)	SOC(e)	SOC(n)		Risk Ra	atio		RR	95%-CI	Weight
RCT Labhardt 2018 MacPherson 2014 Overall effect Heterogeneity: <i>1</i> ² = 92% [72%	134 181 ; 98%], τ ² = 0.	137 490 1797, <i>p</i> < 0.0	44 63	137 278			-	++>	3.05 1.63 2.23	[2.38; 3.89] [1.27; 2.08] [1.21; 4.11]	19.0% 19.0% 38.0%
Cohort Oladele 2018 Reif 2017 Tun 2019 Overall effect Heterogeneity: <i>I</i> ² = 98% [97%	4000 50 256 ; 99%], τ ² = 0.	5789 50 309 1187, <i>p</i> < 0.0	154 305 181	481 330 308		+	+=+		2.16 1.08 1.41 1.48	[1.89; 2.46] [1.05; 1.12] [1.27; 1.57] [1.00; 2.20]	20.4% 21.0% 20.6% 62.0%
Overall effect Heterogeneity: $I^2 = 98\%$ [97% Test for subgroup differences:	; 99%], $\tau^2 = 0$. $\chi_1^2 = 1.21$, df	1504, <i>p</i> < 0.0 = 1 (<i>p</i> = 0.27)	1		0.5	1	2	3.5	1.73	[1.22; 2.45]	100.0%
					Favours So	C	Favours	СОМ			

Fig 3. ART uptake: Community ART initiation versus standard of care among HIV-positive individuals, by study design. COM, community ART initiation; e, number of events; n, number of participants; RCT, randomized controlled trial; RR, risk ratio; SOC, standard of care. MacPherson 2014 cluster-adjusted effect estimate based on the Cochrane method of adjusting for the design effect.

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ART was offered in the community (Fig 3; RR 1.73, 95% CI 1.22 to 2.45; I^2 98%), which translated to an absolute risk difference of 30% (95% CI 10% to 50%) (S1 Fig).

When subgrouped by study design, a stronger effect was seen in RCTs (RR 2.23, 95% CI 1.21 to 4.11) as compared to cohort studies (RR 1.48, 95% CI 1.00 to 2.20); this difference accounted for some of the heterogeneity seen in the overall analysis but not all. There was greater heterogeneity of effect estimates among cohort studies compared to RCTs: 2 cohort studies, one conducted in FSWs (RR 1.41, 95% CI 1.27 to 1.57) and another in adolescents (RR 1.08, 95% CI 1.05 to 1.12), showed smaller differences in ART uptake compared to studies conducted in general HIV endemic communities (RR 2.20, 95% CI 1.56 to 3.12) (S2 Fig). Home ART initiation (explored only in RCTs) had comparable ART uptake (RR 2.23, 95% CI 1.21 to 4.11) to ART offered at other venues in the community (RR 1.48, 95% CI 1.00 to 2.20) (S3 Fig). Same-day ART initiation (RR 1.91, 95% CI 1.05 to 3.46) showed similar ART uptake to initiation within 7 days (RR 1.45, 95% CI 1.30 to 1.62) (S4 Fig).

Retention in care

Four studies compared retention in care between community ART initiation and facility ART initiation and maintenance (standard of care) among HIV-positive individuals at 6–12 months [23,26,28,30]. Retention was higher in the community ART initiation group compared to the facility ART initiation group (RR 1.43, 95% CI 1.32 to 1.54), which translated to a risk difference of 19% (95% CI 11% to 28%) (Figs 4 and §5).

When compared to standard of care (facility-based ART initiation and refills), retention did not appear to differ by population type, ART initiation site, refill site, or frequency of ART refill (Fig 5). One study additionally reported retention among HIV-positive individuals at 24 months and found no difference between community ART initiation and facility ART initiation and facility ART initiation and maintenance (RD 5%, 95% CI –16% to 16%, p = 0.380) at this time point [33]. In pooled analyses restricted to those who initiated ART, retention was no different between

Study	Timepoint	COM(e)	COM(n)	SOC(e)	SOC(n)		Risk Ratio	D		RR	95%-CI	Weight
RCT							I I					
Labhardt 2018	11-14 months	87	137	66	137					1.32	[1.06; 1.63]	13.9%
MacPherson 2014	6 months	129	490	57	278			_		1.21	[0.77; 1.91]	3.1%
Overall effect							-			1.30	[1.07; 1.58]	17.0%
Heterogeneity: $I^2 = 0\%$, $\tau^2 = 0$, p = 0.74											
Cohort												
Tun 2019	12 months	254	309	171	308					1.48	[1.32; 1.66]	50.7%
Reif 2017	12 months	43	50	201	330		- <u>-</u> -			1.41	[1.23; 1.63]	32.2%
Overall effect							•			1.45	[1.33; 1.59]	83.0%
Heterogeneity: $I^2 = 0\%$, $\tau^2 = 0$, <i>p</i> = 0.61											
Overall effect							•			1.43	[1.32; 1.54]	100.0%
Heterogeneity: /2 = 0% [0%; 6	9%], $\tau^2 = 0$, $p = 0.69$							1				
Test for subgroup differences:	$\chi_1^2 = 1.08$, df = 1 (p = 0	0.30)				0.5	1	2	3.5			
						Favours SOC	Favours	COM				

Fig 4. ART retention at 6–12 months among HIV-positive individuals. COM, community ART initiation; e, number of events; n, number of participants; RCT, randomized controlled trial; RR, risk ratio; SOC, standard of care. MacPherson 2014 reflects cluster-adjusted numbers based on the Cochrane method of adjusting for the design effect.

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study arms (S6 Fig; RR 1.15, 95% CI 0.99 to 1.33). The single-arm study Ibiloye 2018 [31] reported 73.2% retention in care among key populations of individuals who initiated ART in the community ART arm after 7 months of follow-up.

An additional study that compared 2 community ART initiation arms reported slightly better retention at 12 months when community ART initiation was combined with facility refills (71%) compared to community-based ART refills (60%); this however did not reach statistical significance (RD -12%, 95% CI -23% to 0.3%) [27].

Viral suppression

Three studies compared viral suppression between community ART initiation and facility ART initiation and maintenance (standard of care) arms among HIV-positive individuals at 12 months; this comparison showed better viral suppression in the community ART group (RR 1.31, 95% CI 1.15 to 1.49) (Fig 6). One study additionally reported 24-month outcomes, which showed no difference in viral load suppression in the community ART arm compared to the facility ART arm (RD 3%, 95% CI –9% to 15%, p = 0.28) at this time point [33]. The threshold for viral load suppression ranged from less than 100 copies/mL [23], to 1,000 copies/mL [28]. In one study, viral suppression in the community ART arm was higher when data

Study	Population	Initiation site	Refill site	Refill frequency	у	Risk Ratio		RR	95%-CI	Weight
Labhardt 2018 MacPherson 2014 Tun 2019 Reif 2017	General General FSW Adolescents	Home Home Community Community	Clinic Clinic Community Community	3m 3m ND 1m	-			1.32 1.21 1.48 1.41	[1.06; 1.63] [0.77; 1.91] [1.32; 1.66] [1.23; 1.63]	13.9% 3.1% 50.7% 32.2%
Overall effect Heterogeneity: <i>I</i> ² = 0% [0%; 69	%], $\tau^2 = 0$, $p = 0.69$				0.5 Favours SC	1 2 DC Favours CC	3.5 M	1.43	[1.32; 1.54]	100.0%

Fig 5. ART retention at 6–12 months among HIV-positive individuals, by implementation strategy. COM, community ART initiation; FSW, female sex workers; ND, not determined; RR, risk ratio; SOC, standard of care. MacPherson 2014 reflects cluster-adjusted numbers based on the Cochrane method of adjusting for the design effect.

Study	Timepoint	COM(e)	COM(n)	SOC(e)	SOC(n)	Ris	k Ratio		RR	95%-CI	Weight
RCT						1					
Labhardt 2018	11-14 months	69	137	47	137	1		1	.47	[1.10; 1.95]	15.3%
Barnabas 2020	12 months	306	427	269	446			1	.19	[1.08; 1.31]	47.2%
Overall effect			564		583		-	1	.26	[1.05; 1.53]	62.5%
Heterogeneity: $I^2 = 47\%$, $\tau^2 = 0$	0.0106, <i>p</i> = 0.17									-	
Cohort											
Tun 2019	12 months	215	309	152	308			1	.41	[1.23; 1.61]	37.5%
Overall effect			309		308		-	1	.41	[1.23; 1.61]	37.5%
Heterogeneity: not applicable											
Overall effect			873		891		•	1	.31	[1.15; 1.49]	100.0%
Heterogeneity: /2 = 61% [0%; 3	89%], $\tau^2 = 0.0066$, $\rho = 0$	0.08									
Test for subgroup differences:	$\chi_1^2 = 0.86$, df = 1 ($p = 0$.35)				0.5 1	2	3.5			
						Favours SOC E	avours COM				

Fig 6. Viral suppression among HIV-positive individuals at 12 months: Community ART initiation versus facility-based care. COM, community ART initiation; e, number of events; n, number of participants; RCT, randomized controlled trial; RR, risk ratio; SOC, standard of care.

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were restricted to men from a South African subgroup (RR 1.39, 95% CI 1.17 to 1.66) [25]. Meta-analysis of viral suppression restricted to those who initiated ART showed no difference in viral suppression between treatment arms at 12 months (RR 1.05, 95% CI 0.86 to 1.29) (S7 Fig).

Two studies compared community ART initiation with hybrid community–facility ART strategies, including either facility-based ART initiation or facility-based ART refill (Table 5). One compared community ART initiation and refills versus facility ART initiation with community ART refills [24,25] and found no difference in viral suppression between these 2 strategies. When analysis was restricted to the South African male subgroup, community ART initiation and maintenance appeared to have better viral suppression than if ART was initiated at the facility and maintained in the community (RR 1.26, 95% CI 1.04 to 1.51) in this study. Another study compared 2 community ART initiation strategies, one with ART refills delivered in the community versus another with refills collected at the health facility; this comparison showed no difference in viral suppression at 12 months between refill strategy arms (RD -7%, 95% CI -20% to 6%) [27].

ART adherence

There was no difference in ART adherence among ART initiators at 6 months in the 2 studies contributing to this comparative analysis (RR 0.92, 95% CI 0.84 to 1.02) (Fig 7). Adherence was assessed as not missing a single dose in the past 4 days [26] or not missing a dose in the past 7 days as assessed by self-report [28].

Table 5. Viral suppression among HIV-positive individuals at 12 months: Community ART initiation and refill versus community–facility hybrid initiation and refill strategies.

Study	Community ART		Hybrid community-faci	lity ART	Effect estimate, 95% CI
	ART initiation	ART refill site	ART initiation	ART refill site	
Barnabas 2020	Community	Community	Facility	Community	RR 1.08, 95% CI 0.98 to 1.19
Amstutz 2021	Community	Community	Community	Facility	RD –7%, 95% CI –20% to 6%

RD, risk difference; RR, risk ratio.

Study	Timepoint	COM(e)	COM(n)	SOC(e)	SOC(r	ו)		Risk Ratio)		RR	95	5%-CI	Weight
Tun 2019 MacPherson 2014	6 months 6 months	177 145	254 181	127 57	171 63		-E	•			0.94 0.90	[0.83; [0.76;	1.06] 1.06]	66.5% 33.5%
Overall effect Heterogeneity: $I^2 = 0\%$, $\tau^2 = 0$, <i>p</i> = 0.65					0.5		1	2	3.5	0.92	[0.84;	1.02]	100.0%
						Favou	irs SOC	Favours	СОМ	0.0				

Fig 7. ART adherence among ART initiators at 6 months. COM, community ART initiation; e, number of events; n, number of participants; RR, risk ratio; SOC, standard of care. MacPherson 2014 reflects cluster-adjusted estimates based on the Cochrane method of adjusting for the design effect.

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Mortality

Overall, there were few events contributing to this outcome. Three studies contributed to the comparative analysis of mortality (at 6 to 12 months), showing no difference in mortality among those who initiated ART in the community compared with those who initiated in the health facility (Fig 8; RR 2.37, 95% CI 0.56 to 10.05). The Ibiloye 2018 study reported overall mortality in its community ART non-comparative cohort study at 3, 6, and 9 months on ART as 3.4%, 3.7%, and 3.9%, respectively. Additionally, Amstutz et al. reported 5% (6/118) mortality in the community ART initiation and refill arm, compared to 0% (0/139) mortality in the hybrid community ART initiation and facility refill arm [27].

Adverse events and social harms

There was variable adverse event reporting, including mild, serious, and severe adverse events, social harms, and opportunistic infection incidence (Table 6). Severe adverse events in community ART initiation arms ranged from 1% to 6% and in facility ART initiation arms ranged from 1% to 2% [23,25,30,33]. There were very few opportunistic infections or social harms; this was however infrequently reported in the included studies.

Certainty of review findings (GRADE assessment)

The certainty of the evidence (a combined assessment of strength of association, methodological quality, heterogeneity, and external validity) for the pooled (RCT and cohort) data on



Fig 8. Mortality among HIV-positive individuals at 12 months. COM, community ART initiation; e, number of events; n, number of participants; RCT, randomized controlled trial; RR, risk ratio; SOC, standard of care.

Study	Adverse events		Opportunistic info	ections	Social harms			
	Community ART initiation	Facility ART initiation	Community ART initiation	Facility ART initiation	Community ART initiation	Facility ART initiation		
Labhardt 2018	6 (4%) events—2 rash, 1 nausea, 1 dizziness, 1 gynecomastia, 1 elevated alanine aminotransferase level	2 events (1%)—2 rash	0 TB cases	2 (1%) TB cases	-	—		
Reif 2017	_	-	_	_	0—increased stigma or unintended disclosure	0—increased stigma or unintended disclosure		
Barnabas 2020 (a)	7 (1%) severe and 7 (1%) serious adverse events	8 (2%) severe and 2 (0.4%) serious adverse events	_	_	2 events	0 events		
Barnabas 2020 (b)	_	5 (1%) severe and 4 (1%) serious adverse events	_	_	_	0 events		
Amstutz 2021 (a)	7 (6%) events—serious adverse events	-	-	-	_	—		
Amstutz 2021 (b)	7 (3%) events—serious adverse events	-	_	-	-	_		

Table 6. Adverse events among HIV-positive.

Barnabas 2020 (a): community ART initiation arm combined with community ART refills. Barnabas 2020 (b): community ART initiation arm combined with facility ART refills. Amstutz 2021 (a): community ART initiation combined with community ART refills. Amstutz 2021 (b): community ART initiation combined with facility ART refills.

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primary outcomes of uptake, retention, and viral suppression among HIV-positive individuals was graded as low to moderate (Table 7); effect estimates were downgraded due to high risk of bias in the contributing observational studies (Table 7). Pooled estimates for adherence were similarly graded as low certainty evidence, due to the inclusion of self-reported outcomes and poor methodological quality. Very few events contributed to the mortality analysis, resulting in very low certainty evidence for this outcome.

Implementation outcomes

Few studies reported on implementation outcomes of community ART initiation, 2 studies reported on cost, and no studies reported on acceptability, penetration, adoption, fidelity, adaptations, feasibility, or sustainability related to community ART initiation.

Of the 2 studies reporting on costs, one reported on the community ART initiation arm only [26], and the second reported cost comparisons of facility ART initiation and community ART initiation across 3 study settings [25]. The Barnabas 2020 study demonstrated some variability across settings, with the cost per person virally suppressed higher with community-based ART initiation compared to facility ART initiation in 2 of the settings (Table 8).

Ongoing studies

We identified 3 ongoing studies being conducted in Indonesia [38], Zimbabwe [39], and Puerto Rico [40]. One study includes adolescents, and 2 studies include key population groups (S3 Table).

Discussion

In this systematic review, we found that making ART initiation available in the community led to increases in ART uptake, better retention, and improved viral suppression (over the course

Certainty assessment							Number of patients		Effect	Certainty	
Number of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other	Community ART initiation	Facility ART initiation	estimate (95% CI)		
ART initiation among PLWH											
5	RCTs and observational	Serious ^a	Serious ^b	Not serious	Not serious	None	4,621/6,466 (71.5%)	747/1,534 (48.7%)	RR 1.73 (1.22 to 2.45)	⊕⊕○○ LOW	
2	RCTs	Not serious	Serious ^b	Not serious	Serious ^c	None	315/503 (62.6%)	107/415 (25.8%)	RR 2.23 (1.21 to 4.11)	⊕⊕○○ LOW	
3	Observational	Serious ^a	Serious ^b	Not serious	Not serious	None	4,306/6,148 (70.0%)	640/1,119 (57.2%)	RR 1.48 (1.00 to 2.20)	⊕000 VERY LOW	
Retention in care among PLWH at 6-12 months											
4	RCTs and observational	Serious ^a	Not serious	Not serious	Not serious	None	513/716 (71.6%)	459/993 (46.2%)	RR 1.44 (1.33 to 1.56)	⊕⊕⊕○ MODERATE	
2	RCTs	Not serious	Not serious	Not serious	Serious ^c	None	216/627 (34.4%)	123/415 (29.6%)	RR 1.30 (1.07 to 1.58)	⊕⊕⊕○ MODERATE	
2	Observational	Serious ^a	Not serious	Not serious	Not serious	None	297/359 (82.7%)	372/638 (58.3%)	RR 1.45 (1.33 to 1.59)	⊕⊕©○ LOW	
Retention in	care among PLWI	H at 24 mor	iths				·				
1	RCT	Not serious	Not serious	Not serious	Very serious ^c	None	88/137 (64%)	81/137(59%)	RD 5% (-16% to 16%)	⊕⊕⊕○ LOW	
Viral suppression among PLWH at 12 months											
3	RCTs and observational	Serious ^a	Not serious	Not serious	Not serious	None	590/873 (67.6%)	468/891 (52.5%)	RR 1.31 (1.15 to 1.49)	⊕⊕⊕○ MODERATE	
2	RCTs	Not serious	Not serious	Not serious	Not serious	None	375/564 (66.5%)	316/583 (54.2%)	RR 1.26 (1.05 to 1.53)	⊕⊕⊕⊕ HIGH	
1	Observational	Serious ^a	Not serious	Not serious	Not serious	None	215/309 (69.6%)	152/308 (49.4%)	RR 1.41 (1.23 to 1.61)	⊕000 VERY LOW	
Viral suppre	ession among PLW	H at 24 mo	nths								
1	RCT	Not serious	Not serious	Not serious	Very serious ^c	None	78/137 (56.9%)	74/137 (54.0%)	RD 3% (-9% to 15%)	⊕000 VERY LOW	
Adherence a	mong ART initiate	es at 6 mont	hs								
2	RCTs and observational	Serious ^a	Not serious	Not serious	Serious ^c	None	322/378 (85.2%)	184/212 (86.8%)	RR 0.92 (0.84 to 1.02)	⊕⊕○○ LOW	
Mortality among PLWH at 6-12 months											
3	RCTs and observational	Serious ^a	Not serious	Not serious	Very serious ^c	None	7/664 (1.1%)	1/572 (0.1%)	RR 2.37 (0.56 to 10.02)	⊕000 VERY LOW	

Table 7. Review evidence certainty assessment (GRADE): Community ART initiation versus facility ART initiation.

PLWH, people living with HIV; RCT, randomized cont rolled trial; RD, risk difference; RR, risk ratio.

^aIncluded observational studies that have methodological concerns—comparison groups drawn from different populations and baseline imbalances were not accounted for in the analysis.

 ${}^{b}I^{2}$ statistic >90%

^cFew studies and events

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of 1 year after ART offer), compared to traditional facility-based ART initiation. Models offering ART initiation in the community increased uptake by 30% and 1-year retention by 19% compared to initiation and maintenance at a traditional health facility. This finding of higher ART uptake in the community ART arm was consistent across study designs and various implementation methods. In one head-to-head comparison of alternative ART refill distribution strategies after community ART initiation, 1-year retention and viral suppression was comparable between community ART refill and facility refill. Another study measured 2-year

Study	Currency	Total cost components	Cost measures reported	Facility ART initiation	Community ART initiation
MacPherson 2014 (Malawi)	2012 US dollars	Community ART initiation (60.3%), staff training (0.6%), community sensitization (0.5%), drugs (3%), consumables (13.8%),	Average cost per participant assessed	—	\$97
		equipment (8.1%), other recurrent items (13.7%); excludes HIV testing costs	Average cost per participant initiated on ART	_	\$127
Barnabas 2020 (South Africa ^a)	2018 US dollars	Cost of ART, trimethoprim-sulfamethoxazole, laboratory testing, personnel, supplies, fuel, and overheads	Annual cost of community-based ART per client	\$249	\$312
			Annual cost per person virally suppressed	\$422	\$452
Barnabas 2020 (South Africa ^b)	2018 US dollars	Cost of ART, trimethoprim-sulfamethoxazole, laboratory testing, personnel, supplies, fuel, and overheads	Annual cost of community-based ART per client	\$249	\$308
			Annual cost per person virally suppressed	\$402	\$380
Barnabas 2020 (Uganda)	2018 US dollars	Cost of ART, trimethoprim-sulfamethoxazole, laboratory testing, personnel, supplies, fuel, and overheads	Annual cost of community-based ART per client	\$163	\$217
			Annual cost per person virally suppressed	\$214	\$275

Table 8. Cost analyses reported in included studies.

^aMidlands Kwazulu-Natal, South Africa. ^bNorthern Kwazulu-Natal, South Africa.

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outcomes and found viral suppression and retention to be no different at this time point between those offered ART initiation in the community and those offered facility initiation. There were too few events to confidently determine the effect of community ART initiation on adherence and mortality. Similarly, few studies reported on adverse events or social harms. These findings were based on the synthesis of 8 studies conducted primarily in low- and middle-income African countries with high to moderate HIV burden, representing diversity in geographical locations and population groups in these settings.

The effect of community-based models for initiating ART had consistent effects across implementation strategies and population subgroups. Community ART was provided through a variety of distribution models: ART was provided at home, in mobile vans and tents, or at community locations; ART was initiated by large multidisciplinary teams in some studies, and in others was primarily nurse driven; and ART refills were subsequently distributed either in the community or at health facilities. In comparative analyses, community ART initiation was explored in general HIV endemic communities, men, adolescents, and FSWs, and in one study was seen to be particularly beneficial for men—suggesting that this may be an additional strategy for improving engagement in HIV services for men, who do not routinely attend health services [8,41]. Among key populations, ART uptake was modest, with the lowest uptake reported in PWID; however, comparative analyses also showed improved treatment outcomes for these groups compared to facility-based offer of ART. Future studies focused on population subgroups could help clarify which implementation strategies are most effective, in which settings, and for whom.

Community ART initiation has the potential to expand differentiated service delivery models and move the entire HIV care cascade into the community, a service delivery approach that both addresses structural barriers to attending health services for people living with HIV (PLWH) and is highly relevant during the COVID-19 pandemic, when decongesting health services has become critical—widespread scale-up, however, needs careful consideration. Although the studies in this review were relatively pragmatic in their design, implementation by external partners, additional resources, technical and logistical assistance, and training provided by research teams may mean that more modest outcomes will be seen when these strategies are incorporated into large-scale public health programs with more limited resources [7]. Lessons can be learned from the implementation of community ART models for "stable" HIVpositive people, which demonstrated effectiveness in trials, but when brought to scale highlighted some of the challenges of bringing services into the community, including difficulties with maintaining the ART supply chains, inadequate resources to support communitybased staff, patient concerns regarding HIV-related stigma in the community, and patient preferences for facility-based care in some instances [42–45].

The mechanism by which offering ART in the community improves retention and viral suppression appears to be through greater ART uptake and reduced loss to follow-up prior to ART initiation. Once ART has been initiated, outcomes are similar to those of individuals who initiate ART at a health facility. Analyses of the effect of offering ART to all PLWH on retention and viral suppression compared to analyses that were restricted to only those who initiated ART showed remarkably improved retention and viral suppression at 1 year in the former compared to smaller differences between community and facility arms in the latter. This suggests that expanding the number of PLWH who initiate ART by offering initiation in the community could have a substantial impact on reaching ART coverage goals, if these effects can be reproduced at scale [46].

Few studies included in this review assessed implementation outcomes beyond cost. Additional study findings regarding fidelity to intervention protocols, challenges and required adaptations, explorations of variability in PLWH and provider preferences and acceptability, provider and health system adoption, and sustainability could aid future implementation and should be incorporated into future study design and reporting [47–50].

Our review findings were limited by there being few studies for inclusion, the incorporation of observational data of poor methodological quality, and short-term HIV treatment outcome measures. For systematic reviews evaluating implementation strategies, the inclusion of observational and programmatic data is critical; however, in order to generate robust and relevant synthesized results, high-quality evidence is needed. Assessments of the methodological quality of observational studies included in this review showed observational studies to be of low quality according to risk of bias tools; this was in large part due to the inclusion of data with observed baseline imbalances in participant characteristics between intervention and comparison groups. Although some studies conducted analyses (e.g., interrupted time series) to adjust for selection, these model outputs could not be included in meta-analyses, and therefore raw unadjusted data were pooled with RCT data [29]. Results from observational studies were however consistent with RCT effects, with no differences between study design subgroup estimates, supporting the pooling of these results. The majority of studies reported HIV treatment outcomes at 1 year or less, with the exception of one study, where 2-year viral suppression showed more moderate treatment outcomes as compared to outcomes at 1 year [33]; it is therefore difficult to draw conclusions on the long-term outcomes of offering ART initiation in the community.

Based on data from a limited set of studies, community ART initiation appears to increase ART uptake and as a result shows better viral suppression and retention in care compared to facility ART initiation and refill. Future research should explore which community ART initiation and refill models are most effective for specific populations, evaluate strategies outside of the African context, and report on long-term and implementation outcomes, to facilitate the incorporation of these strategies into HIV programs.

Supporting information

S1 Appendix. PubMed search strategy. (DOCX)

S2 Appendix. Detailed risk of bias assessments. (DOCX)

S1 Fig. ART uptake among HIV-positive individuals: Risk difference. (PNG)

S2 Fig. ART uptake among HIV-positive individuals, by population group. (PNG)

S3 Fig. ART uptake among HIV-positive individuals, by community ART initiation location.

(PNG)

S4 Fig. ART uptake among HIV-positive individuals, by time to ART initiation. (PNG)

S5 Fig. Retention in care among HIV-positive individuals: Risk difference. (PNG)

S6 Fig. Retention in care among ART initiators. (PNG)

S7 Fig. Viral suppression among ART initiators. (PNG)

S1 Table. PRISMA checklist. (DOC)

S2 Table. Detailed PRECIS-2 rating. (DOCX)

S3 Table. Ongoing studies. (DOCX)

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References

- World Health Organization. Guidelines for managing advanced HIV disease and rapid initiation of antiretroviral therapy. Geneva: World Health Organization; 2017 [cited 2018 Aug 11]. Available from: http://apps.who.int/iris/bitstream/handle/10665/255884/9789241550062-eng.pdf?sequence=1.
- Rosen S, Fox MP. Retention in HIV care between testing and treatment in sub-Saharan Africa: a systematic review. PLoS Med. 2011; 8(7):e1001056. https://doi.org/10.1371/journal.pmed.1001056 PMID: 21811403
- Grimsrud A, Barnabas RV, Ehrenkranz P, Ford N. Evidence for scale up: the differentiated care research agenda. J Int AIDS Soc. 2017; 20(Suppl 4):22024. https://doi.org/10.7448/IAS.20.5.22024 PMID: 28770588
- Sharma M, Ying R, Tarr G, Barnabas R. Systematic review and meta-analysis of community and facility-based HIV testing to address linkage to care gaps in sub-Saharan Africa. Nature. 2015; 528(7580): S77–85. https://doi.org/10.1038/nature16044 PMID: 26633769
- Suthar AB, Ford N, Bachanas PJ, Wong VJ, Rajan JS, Saltzman AK, et al. Towards universal voluntary HIV testing and counselling: a systematic review and meta-analysis of community-based approaches. PLoS Med. 2013; 10(8):e1001496. https://doi.org/10.1371/journal.pmed.1001496 PMID: 23966838
- International AIDS Society. Differentiated service delivery. Geneva: International AIDS Society; 2018 [cited 2018 Sep 18]. Available from: http://www.differentiatedservicedelivery.org.
- Long L, Kuchukhidze S, Pascoe S, Nichols BE, Fox MP, Cele R, et al. Retention in care and viral suppression in differentiated service delivery models for HIV treatment delivery in sub-Saharan Africa: a rapid systematic review. J Int AIDS Soc. 2020; 23(11):e25640. https://doi.org/10.1002/jia2.25640 PMID: 33247517
- Eshun-Wilson I, Rohwer A, Hendricks L, Oliver S, Garner P. Being HIV positive and staying on antiretroviral therapy in Africa: a qualitative systematic review and theoretical model. PLoS ONE. 2019; 14(1): e0210408. https://doi.org/10.1371/journal.pone.0210408 PMID: 30629648
- Moher D, Liberati A, Tetzlaff J, Altman DG, PRISMA Group. Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement. PLoS Med. 2009; 6(7):e1000097. https://doi.org/ 10.1371/journal.pmed.1000097 PMID: 19621072
- Awotiwon A, Amankwaa S, Young T, Eshun-Wilson I, Geng E. Community-based ART initiation for uptake and retention in care in HIV positive individuals. PROSPERO CRD42019130272. National Institute for Health Research; 2019 [cited 2021 May 20]. Available from: https://www.crd.york.ac.uk/ PROSPERO/display_record.php?RecordID=130272.
- 11. World Health Organization. Consolidated guidelines on the use of antiretroviral drugs for treating and preventing HIV infection. Geneva: World Health Organization; 2013 [cited 2021 May 20]. Available from: http://www.who.int/hiv/pub/guidelines/arv2013/download/en/.
- Kredo T, Ford N, Adeniyi FB, Garner P. Decentralising HIV treatment in lower- and middle-income countries. Cochrane Database Syst Rev. 2013; 6:CD009987. https://doi.org/10.1002/14651858. CD009987.pub2 PMID: 23807693
- Higgins JP, Altman DG, Gotzsche PC, Juni P, Moher D, Oxman AD, et al. The Cochrane Collaboration's tool for assessing risk of bias in randomised trials. BMJ. 2011; 343:d5928. https://doi.org/10.1136/ bmj.d5928 PMID: 22008217

- Wells GA, Shea B, O'Connell D, Peterson J, Welch V, Losos M, et al. The Newcastle-Ottawa Scale (NOS) for assessing the quality if nonrandomized studies in meta-analyses. Ottawa: Ottawa Hospital Research Institute; 2018 [2018 Sep 14]. Available from: <u>http://www.ohri.ca/programs/clinical_epidemiology/oxford.asp</u>.
- Loudon K, Treweek S, Sullivan F, Donnan P, Thorpe KE, Zwarenstein M. The PRECIS-2 tool: designing trials that are fit for purpose. BMJ. 2015; 350:h2147. https://doi.org/10.1136/bmj.h2147 PMID: 25956159
- Proctor EK, Powell BJ, McMillen JC. Implementation strategies: recommendations for specifying and reporting. Implement Sci. 2013; 8:139. https://doi.org/10.1186/1748-5908-8-139 PMID: 24289295
- Veroniki AA, Jackson D, Viechtbauer W, Bender R, Bowden J, Knapp G, et al. Methods to estimate the between-study variance and its uncertainty in meta-analysis. Res Synth Methods. 2016; 7(1):55–79. https://doi.org/10.1002/irsm.1164 PMID: 26332144
- R Core Team. R: a language and environment for statistical computing. Vienna: R Foundation for Statistical Computing; 20203 RStudio Version 1.2.5042.
- Guyatt GH, Oxman AD, Kunz R, Brozek J, Alonso-Coello P, Rind D, et al. GRADE guidelines 6. Rating the quality of evidence—imprecision. J Clin Epidemiol. 2011; 64(12):1283–93. https://doi.org/10.1016/j. jclinepi.2011.01.012 PMID: 21839614
- Guyatt GH, Oxman AD, Kunz R, Woodcock J, Brozek J, Helfand M, et al. GRADE guidelines: 8. Rating the quality of evidence—indirectness. J Clin Epidemiol. 2011; 64(12):1303–10. https://doi.org/10.1016/ j.jclinepi.2011.04.014 PMID: 21802903
- Guyatt GH, Oxman AD, Montori V, Vist G, Kunz R, Brozek J, et al. GRADE guidelines: 5. Rating the quality of evidence—publication bias. J Clin Epidemiol. 2011; 64(12):1277–82. https://doi.org/10.1016/j. jclinepi.2011.01.011 PMID: 21802904
- Guyatt GH, Oxman AD, Vist G, Kunz R, Brozek J, Alonso-Coello P, et al. GRADE guidelines: 4. Rating the quality of evidence—study limitations (risk of bias). J Clin Epidemiol. 2011; 64(4):407–15. https://doi.org/10.1016/j.jclinepi.2010.07.017 PMID: 21247734
- 23. Labhardt ND, Ringera I, Lejone TI, Klimkait T, Muhairwe J, Amstutz A, et al. Effect of offering same-day ART vs usual health facility referral during home-based HIV testing on linkage to care and viral suppression among adults with HIV in Lesotho: the CASCADE randomized clinical trial. JAMA. 2018; 319 (11):1103–12. https://doi.org/10.1001/jama.2018.1818 PMID: 29509839
- 24. Barnabas R, van Rooyen H, Asiimwe S, van Heerden A, Pillay D, Szpiro A, et al. Community ART increases viral suppression and eliminates disparities for African men. Conference on Retroviruses and Opportunistic Infections; 2020 Mar 8–11; Boston, MA, USA.
- Barnabas RV, Szpiro AA, van Rooyen H, Asiimwe S, Pillay D, Ware NC, et al. Community-based antiretroviral therapy versus standard clinic-based services for HIV in South Africa and Uganda (DO ART): a randomised trial. Lancet Glob Health. 2020; 8(10):e1305–15. <u>https://doi.org/10.1016/S2214-109X(20)</u> 30313-2 PMID: 32971053
- MacPherson P, Lalloo DG, Webb EL, Maheswaran H, Choko AT, Makombe SD, et al. Effect of optional home initiation of HIV care following HIV self-testing on antiretroviral therapy initiation among adults in Malawi: a randomized clinical trial. JAMA. 2014; 312(4):372–9. <u>https://doi.org/10.1001/jama.2014.6493</u> PMID: 25038356
- Amstutz A, Lejone TI, Khesa L, Kopo M, Kao M, Muhairwe J, et al. Offering ART Refill Through Community Health Workers Versus Clinic-Based Follow-Up After Home-Based Same-Day ART Initiation: The VIBRA Cluster-Randomised Clinical Trial. Available from SSRN: https://ssrn.com/abstract=3824772 or http://dx.doi.org/10.2139/ssrn.3824772
- Vu L, Tun W, Apicella L, Casalini C, Makyao N, Tsang S, et al. Community-based antiretroviral therapy (ART) delivery for female sex workers in Tanzania: intervention model and baseline findings. AIDS Care. 2020; 32(6):729–34. https://doi.org/10.1080/09540121.2019.1640846 PMID: 31298058
- Oladele EA, Badejo OA, Obanubi C, Okechukwu EF, James E, Owhonda G, et al. Bridging the HIV treatment gap in Nigeria: examining community antiretroviral treatment models. J Int AIDS Soc. 2018; 21(4):e25108. https://doi.org/10.1002/jia2.25108 PMID: 29675995
- **30.** Reif L, Bertrand R, Rivera V, Jospeh B, Anglade B, Pape JW, et al. A novel model of community cohort care for HIV-infected adolescents improves outcomes. Top Antivir Med. 2017; 25(1s):355s–6s.
- Ibiloye O, Decroo T, Eyona N, Eze P, Agada P. Characteristics and early clinical outcomes of key populations attending comprehensive community-based HIV care: experiences from Nasarawa State, Nigeria. PLoS ONE. 2018; 13(12):e0209477. https://doi.org/10.1371/journal.pone.0209477 PMID: 30571744
- Alsallaq RA, Baeten JM, Celum CL, Hughes JP, Abu-Raddad LJ, Barnabas RV, et al. Understanding the potential impact of a combination HIV prevention intervention in a hyper-endemic community. PLoS ONE. 2013; 8(1):e54575. https://doi.org/10.1371/journal.pone.0054575 PMID: 23372738

- 33. Amstutz A, Brown JA, Ringera I, Muhairwe J, Lejone TI, Klimkait T, et al. Engagement in care, viral suppression, drug resistance, and reasons for nonengagement after home-based same-day antiretroviral therapy initiation in Lesotho: a two-year follow-up of the CASCADE Trial. Clin Infect Dis. 2020; 71 (10):2608–14. https://doi.org/10.1093/cid/ciz1126 PMID: 31781759
- Labhardt ND, Ringera I, Lejone TI, Masethothi P, Thaanyane T, Kamele M, et al. Same day ART initiation versus clinic-based pre-ART assessment and counselling for individuals newly tested HIV-positive during community-based HIV testing in rural Lesotho—a randomized controlled trial (CASCADE trial). BMC Public Health. 2016; 16:329. https://doi.org/10.1186/s12889-016-2972-6 PMID: 27080120
- Labhardt ND, Ringera I, Lejone TI, Amstutz A, Klimkait T, Muhairwe J, et al. Effect and cost of two successive home visits to increase HIV testing coverage: a prospective study in Lesotho, Southern Africa. BMC Public Health. 2019; 19(1):1441. https://doi.org/10.1186/s12889-019-7784-z PMID: 31676001
- Tun W, Apicella L, Casalini C, Bikaru D, Mbita G, Jeremiah K, et al. Community-based antiretroviral therapy (ART) delivery for female sex workers in Tanzania: 6-month ART initiation and adherence. AIDS Behav. 2019; 23(Suppl 2):142–52. https://doi.org/10.1007/s10461-019-02549-x PMID: 31197700
- Reif LK, Rivera VR, Bertrand R, Belizaire ME, Joseph JB, Louis B, et al. "FANMI": a promising differentiated model of HIV care for adolescents in Haiti. J Acquir Immune Defic Syndr. 2019; 82(1):e11–3. https://doi.org/10.1097/QAI.00000000002088 PMID: 31107305
- Subronto YW. HIV awal (early) testing & treatment Indonesia project intervention phase. ClinicalTrials. gov. 2018 Sep 6 [cited 2021 Apr 26]. Available from: https://ClinicalTrials.gov/show/NCT03659253.
- Ferrand R. Community based interventions to improve HIV outcomes in youth: a cluster randomised trial in Zimbabwe (CHIEDZA). ClinicalTrails.gov. 2018 Oct 25 [cited 2021 Apr 26]. Available from: https://ClinicalTrials.gov/show/NCT03719521.
- Metsch L, Santana J, De Leon SM, Feaster D, Schackman B, Davila G, et al. Enhanced access to HIV care for drug users in San Juan, Puerto Rico (Proyecto PACTo). ClinicalTrials.gov. 2013 Feb 15 [cited 2021 Apr 26]. Available from: https://clinicaltrials.gov/show/NCT01792752.
- Sharma M, Barnabas RV, Celum C. Community-based strategies to strengthen men's engagement in the HIV care cascade in sub-Saharan Africa. PLoS Med. 2017; 14(4):e1002262. <u>https://doi.org/10. 1371/journal.pmed.1002262</u> PMID: 28399122
- Zakumumpa H, Rujumba J, Kwiringira J, Katureebe C, Spicer N. Understanding implementation barriers in the national scale-up of differentiated ART delivery in Uganda. BMC Health Serv Res. 2020; 20 (1):222. https://doi.org/10.1186/s12913-020-5069-y PMID: 32183796
- Eshun-Wilson I, Mukumbwa-Mwenechanya M, Kim H-Y, Zannolini A, Mwamba CP, Dowdy D, et al. Differentiated care preferences of stable patients on antiretroviral therapy in Zambia: a discrete choice experiment. J Acquir Immune Defic Syndr. 2019; 81(5):540–6. <u>https://doi.org/10.1097/QAI.00000000002070 PMID: 31021988</u>
- Roy M, Bolton Moore C, Sikazwe I, Holmes CB. A review of differentiated service delivery for HIV treatment: effectiveness, mechanisms, targeting, and scale. Curr HIV/AIDS Rep. 2019; 16(4):324–34. https://doi.org/10.1007/s11904-019-00454-5 PMID: 31230342
- Mudavanhu M, West NS, Schwartz SR, Mutunga L, Keyser V, Bassett J, et al. Perceptions of community and clinic-based adherence clubs for patients stable on antiretroviral treatment: a mixed methods study. AIDS Behav. 2020; 24(4):1197–206. https://doi.org/10.1007/s10461-019-02681-8 PMID: 31560093
- 46. Joint United Nations Programme on HIV/AIDS. Prevailing against pandemics: by putting people at the centre. Geneva: Joint United Nations Programme on HIV/AIDS; 2020 [cited 2021 Apr 26]. Available from: https://aidstargets2025.unaids.org/assets/images/prevailing-against-pandemics_en.pdf.
- Proctor EK, Silmere H, Raghavan R, Hovmand P, Aarons GA, Bunger A, et al. Outcomes for implementation research: conceptual distinctions, measurement challenges, and research agenda. Adm Policy Ment Health. 2011; 38:65–76. https://doi.org/10.1007/s10488-010-0319-7 PMID: 20957426
- Eshun-Wilson I, Kim HY, Schwartz S, Conte M, Glidden DV, Geng EH. Exploring relative preferences for HIV service features using discrete choice experiments: a synthetic review. Curr HIV/AIDS Rep. 2020; 17(5):467–77. https://doi.org/10.1007/s11904-020-00520-3 PMID: 32860150
- Lewis CC, Fischer S, Weiner BJ, Stanick C, Kim M, Martinez RG. Outcomes for implementation science: an enhanced systematic review of instruments using evidence-based rating criteria. Implement Sci. 2015; 10(1):155. https://doi.org/10.1186/s13012-015-0342-x PMID: 26537706
- 50. Carroll C, Patterson M, Wood S, Booth A, Rick J, Balain S. A conceptual framework for implementation fidelity. Implement Sci. 2007; 2(1):40. https://doi.org/10.1186/1748-5908-2-40 PMID: 18053122