

Understanding care linkage and engagement across 15 adolescent clinics: provider perspectives and implications for newly HIV-infected youth

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Abstract:

The National HIV/AIDS Strategy emphasizes rapid care linkage and engagement for HIV-infected individuals, though many adolescents are never tested, delay entering care, and frequently drop out. We conducted 183 staff interviews at 15 adolescent medicine clinics (baseline, $n = 64$; Year 1, $n = 60$; Year 2, $n = 59$). We used a constant comparative thematic method to examine how providers approached and discussed care linkage/engagement. Qualitative analyses revealed differences in providers' conceptualizations of linkage and engagement. Providers saw linkage as mechanistic and health system driven. It was defined by number of clinic visits and involved relatively little youth agency. In contrast, providers defined engagement by youths' responsibility and participation in their own care. Linkage and engagement are related but distinct aspects of care that require different resources and levels of staff involvement. Integrating an understanding of these differences into future interventions will allow clinic staff to help youth improve long-term health outcomes.

Keywords: adolescent healthcare | care linkage | HIV | engagement

Article:

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UNDERSTANDING CARE LINKAGE AND ENGAGEMENT ACROSS 15 ADOLESCENT CLINICS: PROVIDER PERSPECTIVES AND IMPLICATIONS FOR NEWLY HIV-INFECTED YOUTH

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The National HIV/AIDS Strategy emphasizes rapid care linkage and engagement for HIV-infected individuals, though many adolescents are never tested, delay entering care, and frequently drop out. We conducted 183 staff interviews at 15 adolescent medicine clinics (baseline, $n = 64$; Year 1, $n = 60$; Year 2, $n = 59$). We used a constant comparative thematic method to examine how providers approached and discussed care linkage/engagement. Qualitative analyses revealed differences in providers' conceptualizations of linkage and engagement. Providers saw linkage as mechanistic and health system driven. It was defined by number of clinic visits and involved relatively little youth agency. In contrast, providers defined engagement by youths' responsibility and participation in their own care. Linkage and engagement are related but distinct aspects of care that require different resources and levels of staff involvement. Integrating an understanding of these differences into future interventions will allow clinic staff to help youth improve long-term health outcomes.

INTRODUCTION

The organization of HIV clinical care for adolescents has changed profoundly during the past decade. HIV-infected adolescents have become a significant proportion of the larger epidemic (Centers for Disease Control and Prevention [CDC], 2013) concurrently with advances in HIV therapeutics that—in the United States—have led clinicians to label HIV as a chronic condition and have allowed for decades of health and well-being. Achieving health and well-being for people living with chronic conditions is contingent upon an accessible and sustainable system of care that is responsive to patients' changing needs over the life course (Oni et al., 2014). The challenges of adapting chronic care models to HIV care, at the individual level, are highlighted by evidence that many HIV-infected adolescents are never tested, often fail timely entry into care, drop out at disproportionately high rates, and transition to adult services in ways that often lead to long interruptions of care (Tanner et al., 2016; Whiteside et al., 2014).

On a population level, the HIV care continuum illustrates the difficulty of living with HIV as a chronic condition (Greenberg, Purcell, Gordon, Barasky, & del Rio, 2015; Mugavero, Amico, Horn, & Thompson, 2013; Whiteside et al., 2014). Stages within the care continuum—testing, care linkage and engagement, and medication initiation—provide sequential markers toward a clinical and public health goal of sustained viral suppression. These markers represent points where surveillance data can demonstrate the adequacy of health care systems in meeting the needs of HIV-infected individuals (Das et al., 2013). For example, evidence that a substantial proportion of newly diagnosed persons fail initial care linkage led to intensive case management and patient navigator interventions to facilitate this key early care continuum outcome (E. M. Gardner, McLees, Steiner, del Rio, & Burman, 2011; L. I. Gardner et al., 2009; Giordano et al., 2007). Likewise, interventions to improve care engagement and antiretroviral adherence are being implemented, although seldom in the context of a comprehensive approach to testing, care linkage, and ultimate care engagement (Mugavero et al., 2013). Existing efforts are largely focused on adults rather than adolescents, who comprise over a quarter of new HIV infections in the U.S. (CDC, 2015); adaptation of the care continuum model to include issues relevant to HIV-infected adolescents is relatively recent (Kurth, Lally, Choko, Inwani, & Fortenberry, 2015).

Adolescents face unique challenges to entry into and maintenance of HIV care with the frequent clinic visits characteristic of contemporary HIV medicine and adherence to daily medication regimens (Zanoni & Mayer, 2014). Structural barriers to care such as underemployment, unstable housing, poverty, substance use, lack of educational opportunities, stigma, and lower levels of family support further constrain adolescents' behaviors (e.g., HIV management and health-promoting decisions) (Philbin, Tanner, DuVal, Ellen, Kapogiannis, et al., 2014). Adolescents test for HIV more frequently in community-based rather than clinic-based venues (Swenson, Hadley, Houck, Dance, & Brown, 2011), which have lower rates of linkage success (CDC, 2011) due to the structural fragmentation of testing and care sites (Tanner et al., 2013). Adolescents also delay care linkage for longer periods and are less likely to enter care than older adults (Castle et al., 2011; Grant et al., 2006; Hall et al., 2012; Hosek, Harper, Lemos, & Martinez, 2008; Minniear et al., 2013). These structural barriers are further complicated by adolescents' unique developmental stage, which impacts coping skills, impulsivity, risk taking, and inattention to details and may complicate medication adherence and care engagement. Appointment adherence is challenging; adolescents miss about one-third of scheduled visits, and approximately 30% drop out of care (Minniear et al., 2013). Adolescents also have lower rates of viral suppression than adults (~6% vs. 29%) and higher rates of virologic rebound (Ryscavage, Anderson, Sutton, Reddy, & Taiwo, 2011). For adolescents, care linkage and engagement more often resemble a series of recurrent passages than a continuum, with care interrupted by periods of disengagement followed by re-established care (Zanoni & Mayer, 2014).

This article addresses the relative lack of data to guide the development of integrated clinical services for HIV-infected adolescents. Improvement of such services is particularly important because adolescence is often ambiguously represented in service delivery systems, divided between pediatric and adult specialties; HIV medicine is also largely a domain of adult specialties, and the way care is often provided (i.e., in a non-youth-friendly way) may affect outcomes (Kasedde, Kapogiannis, McClure, & Luo, 2014; Kendall et al., 2015). Since systems of care for HIV-infected adolescents lack a clear means for assessment of best practices (Kapogiannis, Legins, Chandan, & Lee, 2014), we interviewed health services providers with specific experience in the care and treatment of HIV-infected adolescents. These interviews addressed two key stages on the HIV care continuum: care linkage and engagement. Understanding potential distinctions between these two early stages of care is important to guide program design to reduce adolescent-specific barriers, appropriately direct resources, and provide supports for different types of needs.

METHODS

Data were collected through the evaluation of SMILE (Strategic Multisite Initiative for the Identification, Linkage, and Engagement in Care of Youth with Undiagnosed HIV Infection), a program developed to improve care linkage and engagement for HIV-infected adolescents (Philbin, Tanner, DuVal, Ellen, Xu, et al., 2014). SMILE created formal networks among testing, referral, and clinical agencies (including local health departments), and supported care linkage through the provision of case management and patient navigator services for newly diagnosed adolescents (Tanner et al., 2013). Details of the SMILE intervention, including linkage and engagement outcomes, are described elsewhere (Philbin, Tanner, DuVal, Ellen, Xu, et al., 2014).

The SMILE qualitative evaluation included 183 semistructured individual interviews at 15 geographically dispersed adolescent HIV medicine clinics in the U.S. (e.g., San Francisco, New Orleans, Miami, Baltimore, and Chicago). In addition to speaking with Adolescent Medicine Trials Network for HIV/AIDS Interventions (ATN) staff (i.e., outreach workers and linkage coordinators), we used purposive sampling to conduct interviews with medical and social service providers at each site (e.g., case managers, nurses, social workers, and physicians). These interviews occurred across three annual site visits from 2010 to 2012 (prior to SMILE initiation, $n = 64$; Year 1, $n = 60$; Year 2, $n = 59$). The first or second author conducted interviews; no financial incentives were provided. Analyses of the interviews conducted prior to SMILE initiation and at Year 1 demonstrated the importance of providers' different approaches to linkage and engagement; thus, we added questions to the Year 2 interview guide that explicitly asked participants to discuss their understanding and conceptualization of care linkage and engagement. Interviews were digitally recorded, transcribed, and uploaded into Atlas.ti 6.2 for data management and to assist with coding, analysis, and interpretation. All interviewees provided verbal consent; Institutional Review Boards at the University of North Carolina Greensboro and all participating sites approved the study protocol.

Data were analyzed using a constant comparative method to identify differences and similarities in providers' descriptions and approaches toward care linkage and engagement (Buetow, 2010; Glaser & Strauss, 1967). Two team members independently read and coded each transcript using a line-by-line method to create an initial code dictionary. We then created a list of thematic codes based on the existing literature and integrated it into the initial code dictionary to ensure that both theory-based and emergent concepts were included. Team members then independently cross-coded a random sample of 33% of transcripts to refine the code dictionary, which was subsequently reviewed by other team members (MacQueen, McLellan, Kay, & Milstein, 1998). Next, we created a data table to summarize and refine codes (Glaser & Strauss, 1967) and a site matrix to compare site similarities and variations in descriptions of linkage and engagement. As final coding occurred, the research team conducted a sequence of weekly meetings to develop additional codes and resolve discrepancies. Researchers applied the finalized structure to all transcripts, and there was high consistency between raters. Any discrepancies were resolved through discussion involving the coders and, when needed, the primary investigators.

RESULTS

CARE LINKAGE VERSUS ENGAGEMENT

Respondents positioned the relationship between adolescents and care systems differently for care linkage and care engagement. Care linkage was viewed as a procedure performed on an adolescent; engagement was adapted based on a patient's actions: "when I link something that means I'm actually doing a physical act," whereas an engaged patient is one who "showed up at the clinic on his own" (linkage coordinator; Site C). Linkage was also described as a relatively uniform process regardless of patient characteristics, whereas engagement was a personalized process tailored to a patient's needs. A social worker noted:

Linking is more action oriented, there's a process of steps in making things happen, and engagement really has to do a lot with relationship building and figuring out the needs of each patient and being able to do that over a period of time. (Site L)

While clinic staff worked to support youth and to be advocates, linkage was primarily about “getting kids to an appointment” (outreach worker; Site P), providing information, and arranging blood draws. There was often little patient-client interaction prior to linkage, though some social service providers described a “touch” rate where they tried to send text messages before the first clinic visit. For some respondents, care linkage was described as a countermeasure for what they attributed to a patient's perceived incapacity: “Hey, I need some kind of assistance, because I'm not going to access this. I'm still in denial” (linkage coordinator; Site D).

In contrast to linkage, which was defined by an action (i.e., a clinic visit or lab tests), practitioners defined engagement as youth taking responsibility and actively participating in their own care, showing up without prodding, and accepting their diagnosis. In addition, staff often depicted engagement as a process that involved not only presenting at the clinic, but as “having it together” (outreach worker; Site C) more broadly in terms of housing, employment, and self-care. A few providers supplemented this definition by noting that engagement was only achieved once an appropriate clinical plan existed and the adolescents had “actually gotten the clinical services that they need” (outreach worker; Site P).

DIFFERENCES IN DURATION

While descriptions of linkage were typically precise and short, engagement often led to lengthy discussions to articulate what one nurse summarized as “just messy” (Site E). Many respondents noted ways in which engagement contradicted the assumption of a linear care continuum, “engagement is a continuum of care... engagement is a continuum of linkage, but it's continued over a period of time” (care coordinator; Site M). Indeed, engagement itself was a cycle of recurrent entry, drop out, and re-entry; providers never described youth as “dropping out” of linkage activities.

Participants' perceptions of the time frames for linkage and engagement also differed: linkage was described as a rapid event that should occur within a week of referral; in contrast, engagement was a potentially prolonged process, in which the pace was determined by the adolescent. As a social worker noted:

A lot of young people get linked quickly but I would think engagement would take a couple of years, which sounds kind of crazy, but just to kind of get used to coming to the clinic and dealing with that. (Site K)

ACCEPTANCE OF HIV

An adolescent's acceptance of his or her HIV diagnosis was considered vital because it forced an acknowledgment of the potential challenges to medical linkage and engagement. This involved not only an awareness of social barriers that come with being HIV infected, but also that one's life will continually include interactions with a clinical setting. For example, whereas stigma was not reported as a barrier to care linkage “because in most clinical sites you can just walk the adolescent over” (outreach worker; Site M), providers described stigma as a consistent problem for care engagement. Clinicians described adolescents who were “scared and not ready to deal with it or in denial and they have to hide it from family and friends” (social worker; Site L), which often required offering solutions outside of the clinic until

adolescents were ready to engage in medical care. Respondents frequently discussed the challenges of working with adolescents who struggled to accept their diagnoses:

It's a process, it's not—they're accepting and that's okay. There is grief and there's anger and everything else.... Most of them come. They tell us, "I'm not ready to start medications yet." We tell them, "It's no problem. We can give you a break until you're ready." (psychologist; Site N)

One physician described the acceptance portion of engagement as "an internal process for the patient; engagement in care is a process where the person understands that they need to go to a health care facility for the rest of their life. And sometimes that process doesn't happen immediately" (Site N). Providers talked about adolescents who attended clinic visits for a period of time and then grew tired of the work required to be consistently engaged. Providers struggled with this, and with the chronic nature of HIV infection and its treatment. A staff member stressed how engagement differed from linkage in that "engagement is cyclical. I think it's a whole 'nother animal, you know, the long-term willingness to deal with a chronic illness" (linkage coordinator; Site J).

BUILDING RELATIONSHIPS

Respondents repeatedly stressed the importance of building relationships between adolescents and clinic personnel, whether a physician, receptionist, or security guard, for care engagement. While this could happen in one visit, staff said it frequently took months or even years. A nurse stated that having personal relationships was important, "above all else" (Site J), and an outreach worker noted that "a consistent relationship with one medical provider is more than, well, not more important than getting your labs done, but *as* important as getting your labs done at least" (Site B). Although crucial, adolescent-staff relationships were described as particularly challenging. Many adolescents lacked stable relationships with adults, and previous experiences with health care or social service providers often involved child protective services or social workers. As one care coordinator noted, "To assume that a young person is just going to instantly develop a relationship with a physician is ridiculous" (Site B). Respondents also believed these relationships facilitated a safe space in which adolescents could voice their own needs, ask questions, and share relevant challenges. Indeed, providers also associated engagement with how well they knew an adolescent and whether an adolescent would tell them the truth (e.g., about drug use).

BECOMING A "GOOD" PATIENT

Another aspect of effective engagement was the process through which adolescents became "good" patients who took care of themselves, and who would grow into healthy and successful adults. Staff frequently used words like "knowledgeable," "cooperating," and "not complaining" to describe those who were "good," engaged patients. A case manager described what would ideally occur for a youth to be considered engaged:

They are keeping appointments, taking their medications and not just because someone's telling them to. If they don't, they're calling us to reschedule or to acknowledge that they missed an appointment. They're having productive conversations with people related to their health care. They want to stay healthy and they want to start to educate others around them. (Site M)

Getting youth engaged helped prepare them for transition to adult care; this was particularly important because of a perception that adult HIV care providers were much less tolerant of missed or canceled appointments.

The concept of the “good” patient was also discussed by participants who noted the importance of wanting to stay mentally and physically healthy, being adherent, eliminating the behaviors that put the adolescent at risk for HIV (e.g., not using condoms or avoiding sex during treatment for a sexually transmitted infection [STI]), and actively contributing to care-related decisions. Specifically, this meant knowing the names and doses of medications, understanding viral loads and CD4 cell counts, and asking relevant questions about their condition. One outreach worker summarized this perspective: “Engagement means you’re going to your appointments; you’re involved in your care; you’re asking questions; you’re taking your medicine” (Site I).

INVOLVEMENT IN ANCILLARY SERVICES

Respondents described patient involvement with ancillary services (e.g., vocational, educational, or housing support) as a vital component of care behaviors since the acceptance of an HIV diagnosis often required adolescents to address the social and structural factors that facilitated their HIV acquisition and challenge medical engagement. Respondents believed that participation in ancillary services helped to engage adolescents in the clinic more broadly: “You know, not just them attending that first and second medical appointment but being able to help them get the resources that they need: housing, food stamps, ID, etc.” (outreach worker; Site C). Involvement in these ancillary services was seen as important to help adolescents address the issues that would allow them to fully engage in HIV care, particularly for members of groups who were vulnerable based on race/ethnicity, sexual orientation, or economic status:

A lot of the youth are African American and they’re coming out of the closet and they’re also testing positive.... I mean if you feel like you’re bad, or evil, or wrong, or deserving of the HIV because of your sexuality that’s a huge impact on readiness. You got to work with that with the youth before you can even attempt to begin to address their medical care. (nurse; Site I)

DISCUSSION

These data produced three key findings that can help inform the structure of HIV-related health care services for adolescents. First, care linkage was consistently described as a finite procedure with a specific goal—achieving a first medical appointment—whereas engagement was a process of learning and experience, often marked by periods of less than optimal adherence to clinical and treatment regimens or by complete interruption of engagement followed by re-entry into care. The perception of linkage as a relatively straightforward procedure may reflect recent developments of adolescent linkage-related infrastructure and skill enhancement, as earlier work suggests a fragile and complex process often ending in failure (Fortenberry, Martinez, Rudy, & Monte, 2012; Hosek et al., 2008). This perception may also reflect the fact that it is a stage over which providers have more control: staff can bring an adolescent to a clinic and then linkage is complete; engagement is primarily an adolescent’s responsibility. Even though care linkage was described as relatively

straightforward, studies suggest that only two-thirds of newly diagnosed adolescents are successfully linked (Philbin, Tanner, DuVal, Ellen, Xu, et al., 2014). In addition to personal characteristics, youth are more likely to link if they are stably housed, have health insurance, and are referred to a clinic that is adolescent specific; youth linked more quickly are also more likely to become engaged (Philbin et al., 2016). This demonstrates the importance of creating integrated systems of care to ensure that regardless of where young people test—e.g., clinic, community organization, or health fair—they will be rapidly referred to a clinic where they will feel comfortable (Tanner et al., 2014). Then, once at the clinic, even if a young person does not want to be linked to medical care, social service providers can work with him or her to address other factors that could facilitate linkage and, later, engagement (e.g., housing, education, health insurance).

A second key observation is the need to “become a good patient” for engagement to be deemed successful. This emphasis certainly has practical overtones in that proactive engagement in self-care, adoption of healthier lifestyles, clinic appointment keeping, and therapeutic adherence are relevant to reasonable clinical care objectives such as long-term viral suppression, prevention of HIV disease morbidity, and reduction in HIV transmission risk (Hoffmann & Gallant, 2014). However, “becoming a good patient” suggests the imposition of social and behavioral standards often targeted toward disadvantaged youth and sexual or racial/ethnic minorities. It emphasizes the work patients must do to achieve the expectations that result from HIV being seen as chronic, namely to remain virally suppressed and healthy. Similar themes have emerged in transplantation studies where patients must demonstrate that they are worthy of receiving an organ for which many people are on a waiting list (Cass et al., 2007; Furnham, Ariffin, & McClelland, 2007). With adolescent patients, it also seems suggestive of “transitional paternalism” as an appropriate stance for adults (in this case, health professionals) and adolescents with social, developmental, and legal boundaries on behavioral autonomy (Manson, 2015). Social service and medical providers can facilitate a youth’s transition into a “good patient” by moving away from the provider-centered model that is common in pediatric medicine and developing a collaborative model that includes direct patient input. Such a model allows youth to describe how they envision being a “good patient,” and to design their own treatment goals, realistically assess barriers, and practice skills needed to reach the jointly decided goals.

A third key observation was that care engagement, and, thus, retention, is a process rather than a definitive event. This means that assessments of engagement based on number of clinic visits cannot adequately represent the complex issues faced by adolescents (e.g., social barriers and need for specific ancillary services; Mugavero et al., 2009, 2012, 2013). Engagement is most commonly defined as a second HIV-related visit within a given time period after an initial visit (Fleishman, Yehia, Moore, Korthuis, & Gebo, 2012; Giordano et al., 2007), though details vary to include the number of missed visits, the proportion of kept visits to scheduled visits, the number of 3-month intervals with at least one attended medical visit, the existence of a 6-month gap in care, and whether a person kept two visits separated by at least 90 days (Fleishman et al., 2012; Giordano et al., 2007; Mugavero et al., 2012, 2013). The U.S. National HIV/AIDS Strategy differentiates linkage and engagement only by time and the number of clinic visits (Office of National AIDS

Policy, 2015). The ways our respondents described the engagement process differed from the time lines and straightforward definitions offered by the CDC (2016; see also Mugavero et al., 2012). Engagement, in particular, takes incredible energy and time. It includes overcoming social barriers and developing relationships with clinical staff. Even if adolescents attend sufficient visits to meet the CDC's definition of "engaged," it frequently takes much longer for them to embrace the characteristics that would cause a provider to label them as engaged. This framing of care engagement as a cyclical and multiphase process has been demonstrated by other research on youth living with chronic illness (e.g., diabetes, cystic fibrosis; Lotstein et al., 2013; Sobota et al., 2014). This suggests that providers should place additional weight on building relationships to ensure that youth remain connected to the clinic, even if they are not taking their medication or engaging in care.

Adolescent providers' extended approach to engagement made it almost impossible for them to achieve the benchmarks outlined by the CDC and listed in the U.S. National HIV/AIDS Strategy. This suggests the importance of considering whether there is a need to develop alternative national-level benchmarks that match the realities of adolescents' day-to-day lives in HIV clinics. Future research should work with newly diagnosed youth to explore how they conceptualize linkage and engagement. Then, findings from youth could be integrated into provider-focused research to inform interventions that will help increase rates of linkage and engagement for this vulnerable population.

LIMITATIONS

These data examine how adolescent providers across a wide geographic area understand care linkage and engagement. Geographic variability in the availability of ancillary services and support may facilitate linkage and engagement, and thus how staff came to understand them. Adolescent Trials Network sites are all located in major urban centers where the HIV epidemic is concentrated for youth, and these findings may not be generalizable to other settings in the U.S. While this study relied on the perspectives of staff, future studies could enhance our understanding of the adolescent-specific HIV care continuum by interviewing adolescents to see how they understand and experience differences in care linkage and engagement.

CONCLUSIONS

Linkage and engagement are related but distinct aspects of care that require different resources and provider orientation toward youth. The implementation of seamless linkage and engagement services requires a nuanced understanding of the similarities and differences and what each involves. Successful care linkage and engagement, particularly for adolescents, will be possible only if the system of care is designed in an accessible and sustainable way. These findings can help inform the development of comprehensive clinical services for adolescents living with HIV. Integrating an understanding of these differences into future interventions will allow health care and social service providers to help retain youth in clinics to begin antiretroviral medication, improve their engagement in care, and maintain viral suppression for the long term.

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