

# Reliability, Validity, and Factor Structure of the Hopkins Symptom Checklist-25: Population-Based Study of Persons Living with HIV in Rural Uganda

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**Abstract** Depression and anxiety are highly comorbid among people living with HIV (PLHIV), but few instruments for screening or measurement have been validated for use in sub-Saharan Africa. The objective of this study was to determine the reliability, validity, and factor structure of the 25-item Hopkins Symptom Checklist (HSCL) in a population-based sample of PLHIV in rural Uganda. This study was nested within an ongoing population-based cohort of all residents living in Nyakabare Parish, Mbarara District, Uganda. All participants who identified as HIV-positive by self-report were included in this analysis. We performed parallel analysis on the scale items and estimated the internal consistency of the identified sub-scales using ordinal alpha. To assess construct validity we correlated the sub-scales with related constructs, including subjective well being (happiness), food insecurity, and health status. Of 1814 eligible adults in the population, 158 (8.7%) self-reported being HIV positive. The mean age was 41 years, and 68% were women. Mean HSCL-25 scores were higher among women compared with men (1.71 vs. 1.44;  $t = 3.6$ ,  $P < 0.001$ ). Parallel analysis revealed a three-factor structure that explained 83% of the variance: depression (7 items), anxiety (5 items), and

somatic symptoms (7 items). The ordinal alpha statistics for the sub-scales ranged from 0.83 to 0.91. Depending on the sub-scale, between 27 and 41% of the sample met criteria for caseness. Strong evidence of construct validity was shown in the estimated correlations between sub-scale scores and happiness, food insecurity, and self-reported overall health. The HSCL-25 is a reliable and valid measure of mental health among PLHIV in rural Uganda. In cultural contexts where somatic complaints are commonly elicited when screening for symptoms of depression, it may be undesirable to exclude somatic items from depression symptom checklists administered to PLHIV.

**Keywords** HIV · Depression · Anxiety · Screening · Case-finding · Uganda · Sub-Saharan Africa

## Introduction

Depression, which is one of the leading causes of disability worldwide [1], is highly prevalent among people living with HIV (PLHIV) [2, 3]. In studies conducted in sub-Saharan Africa, nearly one in five PLHIV have been found to have major depressive disorder and nearly one in three PLHIV have clinically significant symptoms of depression [4]. These statistics highlight an important public health problem because depression is associated with reduced HIV treatment adherence and adverse health outcomes among PLHIV, such as HIV transmission risk behavior, poorer physical functioning, and hazardous alcohol use [5–8]. In addition, the presence of anxiety symptoms among PLHIV has also been shown to compromise HIV-related outcomes, although less is known in this regard [9].

Identifying and treating depression among PLHIV not only improves access to care and initiation of treatment but

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also improves treatment adherence and prevents disease progression [5, 6, 10]. Accordingly, there has been considerable interest in the use and application of depression screening tools in care settings where specialist psychiatric care is unavailable [11]. Since both depression and HIV may present with similar somatic or physical symptoms, screening for depression among PLHIV can be a difficult undertaking in population-based studies, especially in sub-Saharan Africa [12, 13]. Moreover, there is need to consider cultural differences when using depression scales developed in Western populations to screen for depression in African populations [11, 14]. Overall, the lack of validated screening tools complicates research focusing on mental health and HIV in sub-Saharan Africa [11].

There are few published studies on the validation of screening instruments for detection of both depression and anxiety among PLHIV. In a recently published systematic review of scale validation studies conducted in sub-Saharan Africa [4], the most frequently studied depression screening scales were the Center for Epidemiologic Studies-Depression (CES-D) and the Patient Health Questionnaire (PHQ). Few studies compared the reliability and/or validity of different depression screening scales; Akena et al. [15] was a notable example in comparing the CES-D and PHQ-9 to the Kessler-10. Several studies have identified a somatic factor, suggesting the importance of careful symptom assessment among PLHIV in the setting of depression and/or anxiety. Depending on illness stage and treatment status, PLHIV may experience somatic symptoms such as fatigue, disturbed sleep, and weight loss that could potentially overlap with the manifestations of depressive disorders [16]. Among studies that examined multi-factor scales, such as the Hopkins Symptom Checklist (HSCL) for Depression, most focused solely on the depression subscales and typically excluded any anxiety subscales from consideration. One exception is the study conducted by Kaaya and colleagues [17], who validated a depression and anxiety screening scale among pregnant women in Tanzania. To address this gap in the literature, we examined the reliability and validity of the HSCL for Depression, a 25-item scale that assesses symptoms of both depression and anxiety, in a population-based sample of PLHIV in rural Uganda.

## Methods

### Ethics Statement

We obtained feedback on the design of our study from a community advisory board, comprised of eight community leaders (four men and four women), including the district community development officer and a person living with

HIV volunteering as a counselor at the local HIV clinic. Ethical approval for all study procedures was obtained from the Partners Human Research Committee, Massachusetts General Hospital; and the Institutional Review Committee, Mbarara University of Science and Technology. Consistent with national guidelines, we received clearance for the study from the Uganda National Council for Science and Technology and from the Research Secretariat in the Office of the President.

### Study Population and Design

The study was conducted in Nyakabare Parish of Mbarara District, located approximately 275 km southwest of Kampala, the capital city of Uganda. The local population largely depends on subsistence agriculture for income and consumption, and both food and water insecurity are common [18, 19]. We used an iterative process, involving conversations with local officials and field investigations, to select prospective study sites. We selected Nyakabare Parish because of its tractable size, both in terms of population and in terms of geographic area, and because the village leadership supported our study's presence.

Approximately 3 months prior to survey administration, we conducted a population census within the parish and enumerated all adults aged 18 years and older, as well as emancipated minors (16–18 years). We excluded persons younger than 18 years of age who were not emancipated minors and adults who did not consider Nyakabare their primary place of residence. We also excluded adults who could not communicate effectively or efficiently with research staff, e.g., due to deafness, mutism, or aphasia; and adults with psychosis, neurological damage, acute intoxication, or an intelligence quotient less than 70 (all of which were determined in the field by non-clinical research staff in consultation with a supervisor).

A research assistant who spoke the local language (Runyankore) approached each potentially eligible study participant at their home or (less frequently and only by request) place of work. The research assistant described the study in detail to each person who expressed potential interest in participation. Written informed consent to participate was obtained. Study participants who could not write were permitted to indicate consent with a thumbprint. Participants were interviewed on a one-on-one basis in a private area, out of earshot from other persons.

### Survey Instrument

The survey was programmed into laptop computers for administration in the field, using the Computer Assisted Survey Information Collection (CASIC) Builder™ software program (West Portal Software Corporation, San

Francisco, Calif.). Survey questions were first written in English, translated into Runyankore, and then back translated into English to verify the fidelity of the translated text. The translation and back-translation was an iterative process involving in-depth consultation and pilot testing with 18 key informants.

All study participants were asked to report their HIV serostatus, with the following response options: HIV-positive, HIV-negative, unknown, or refused. Given that it is rare for persons to identify as HIV-positive when they are actually seronegative [20], study participants were categorized as HIV-positive if they indicated as such on the survey. All other participants were categorized as having unknown serostatus. The analyses reported in this manuscript focus specifically on the sample of study participants who identified themselves as HIV-positive.

The 25-item HSCL was originally derived from the 90-item Symptom Checklist (SCL-90) and has traditionally been administered as a 15-item subscale measuring symptoms of depression and a 10-item subscale measuring symptoms of anxiety [21]. Following the scale development study conducted by Bolton and Ndogoni [22] in rural Uganda, we modified the depression subscale by adding a new item (“don’t care what happens to your health”) and deleting one item (“feeling trapped”). Participants rated the frequency of each symptom in the last seven days on a four-point Likert-type scale, ranging from “not at all” (1) to “very much” (4). The total severity score is then calculated as the mean of the items (range 1–4), with higher scores indicating greater symptom severity. In addition to the survey procedures above, two experienced Ugandan psychiatrists scrutinized the translation to ensure that it preserved concepts and connotations beyond simple semantic equivalence.

We also measured several related constructs, including subjective well-being (happiness), food insecurity, and self-reported overall health. Subjective well-being has a well known inverse correlation with psychological distress [23, 24]. To measure subjective wellbeing, we used a one-item happiness scale: “Considering everything together, how would you say things are in your life? Would you say that you are very happy, fairly happy, or not happy?” In many settings in sub-Saharan Africa, food insecurity is one of the most predominant forms of uncertainty experienced in daily living and has a well known positive correlation with psychological distress [25, 26]. To measure food insecurity, we used the 3-item Household Hunger Scale, an abbreviated form of the 9-item Household Food Insecurity Access Scale, which has been validated in this population [18, 27]. Finally, to measure self-reported overall health, we used a one-item scale [28, 29]: “How is your health in general? Is it very good, good, bad, or very bad?”

## Statistical Analysis

Statistical analyses were conducted using Stata/MP software (version 13.1, StataCorp LP, College Station, Tex.) and R (version 3.3.2, The R Foundation for Statistical Computing, Vienna, Austria). We used parallel analysis to inform our decision about the number of factors to retain [30–32]. We calculated ordinal alphas to assess the internal consistency of the identified factors [33]. We examined item-test correlations, and then re-calculated the ordinal alphas after sequentially deleting each of the items in turn.

The subscale scores were calculated by averaging across the subscale items. As with many large-scale epidemiological studies, we did not have access to a gold standard criterion of mental health (e.g., clinical psychiatric assessments for all members of the population) to assess criterion-related validity. Therefore as a preliminary step to identify caseness for depression and anxiety, we used the conventional cutoff of  $>1.75$  that has been used previously [21, 34–36]. Caseness is a binary variable that indicates a positive screen for depression and/or anxiety, equal to 1 if the subscale score exceeds the cutoff and 0 otherwise. In addition, we relied upon several different assessments of construct validity by correlating the subscales with the previously described variables theorized to be associated with mental health status in this context: subjective well-being (happiness), food insecurity, and self-reported overall health. To estimate correlations we computed Spearman’s rank correlation coefficient, with confidence intervals calculated using Fisher’s  $z$  transform [37, 38]. Test statistics from two-sample  $t$  tests were converted to point-biserial correlation coefficients to facilitate comparisons with the Spearman rank correlation coefficients.

## Results

### Summary Statistics

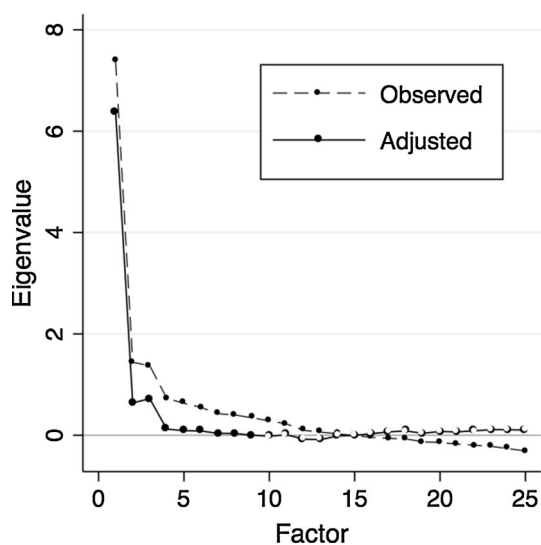
Sociodemographic characteristics of the sample are listed in Table 1. Of the 1814 eligible adults in the population, 158 reported being HIV positive (8.7%). Mean participant age was 41 years (standard deviation [SD], 10.7), 68% were women, and 54% were married or cohabiting. Study participants were evenly distributed among the 8 villages of the parish, with between 17 and 22 PLHIV per village. The mean HSCL-25 score was 1.6 (SD 0.45) and ranged from 1 to 2.8, with mean scores higher among women compared with men (1.71 vs. 1.44;  $t = 3.6$ ,  $P < 0.001$ ).

**Table 1** Summary statistics (N = 154)

	Mean (range) or N (%)
Sex	
Men	52 (33%)
Women	106 (67%)
Age, years	41 (17–70)
Formal education	
None	27 (17%)
Some primary (P1–P6)	67 (42%)
Completed primary (P7–P8)	40 (25%)
More than primary	24 (15%)
Marital status	
Married or cohabitating	86 (54%)
Separated, divorced, or widowed	60 (38%)
Single	11 (7%)

### Factor Structure

Exploratory factor analysis of the 25-item HSCL revealed a dominant factor with an eigenvalue of 7.4 that alone explained 60% of the variance. There were two additional factors both with eigenvalues of 1.4 that explained another 12 and 11% of the variance. Visual examination of the scree confirmed this three-factor structure (Fig. 1). The adjusted eigenvalue was largest for factor 1, followed by steep declines in the adjusted eigenvalues for factors 2 and 3; the remaining adjusted eigenvalues were negligible. Oblique promax rotation yielded a 3-factor structure



**Fig. 1** Scree. The graph shows the observed versus adjusted eigenvalues of the identified factors, in declining order of magnitude. As is shown in the figure, the adjusted eigenvalue was largest for factor 1, followed by steep declines in the adjusted eigenvalues for factors 2 and 3; the remaining adjusted eigenvalues were negligible

solution with a pattern of factor loadings that was easiest to interpret. Factor 1 consisted of 7 items, with factor loadings ranging from 0.49 to 0.76, that were related to depressed mood and negative affect; we labeled this factor the “depression subscale” (Table 2). Factor 2 consisted of 5 items, with factor loadings ranging from 0.46 to 0.84 that were related to agitation or anxiety; we labeled this factor the “anxiety subscale.” Factor 3 consisted of 7 items, with factor loadings ranging from 0.42 to 0.68 that were related to somatic symptoms; we labeled this factor the “somatic symptoms subscale.” The remaining 6 items did not load substantively on any of these identified factors. Correlations between these three factors ranged from 0.4 to 0.5.

### Reliability

The 7-item depression subscale had an ordinal alpha of 0.90. Mean values on this subscale ranged from 1 to 3.7, with a sample mean of 1.51 (SD 0.58); higher scores indicate greater depression symptom severity. Approximately one-quarter of the sample (42 [27%]) met criteria for caseness on this scale. The 5-item anxiety subscale had an ordinal alpha of 0.91, with values ranging from 1 to 3.6 (mean 1.45; SD 0.60); higher scores indicate greater anxiety symptom severity. A similar number of study participants (44 [28%]) met criteria for caseness on the anxiety subscale. The 7-item somatic symptoms subscale had an ordinal alpha of 0.83. Values on the somatic symptoms subscale ranged from 1 to 3.9, with a mean of 1.81 (SD 0.62). A larger proportion of the sample (64 [41%]) met criteria for caseness on the somatic symptoms subscale. Of the participants who met criteria for caseness on the somatic symptoms subscale, only 26 (41%) also met criteria for caseness on the depression subscale; similarly, of the participants who met criteria for caseness on the depression subscale, 16 (38%) did not also meet criteria for caseness on the somatic symptoms subscale. More than one-half of the sample (89 [56%]) met criteria for caseness on at least one subscale, while only 20 (13%) met criteria for caseness on all three subscales.

### Validity

The 7-item depression subscale identified in this study had a statistically significant positive correlation with the modified 15-item depression subscale developed by Bolton & Ngodoni [22] ( $r = 0.87$ ; 95% CI 0.83–0.91). When participants were categorized according to caseness on both subscales, most participants were consistently classified on both subscales (125 [81%]). Twenty-one participants (14%) were classified as having probable depression on the Bolton & Ngodoni subscale but not on the new 7-item depression subscale, while only 8 participants (5%)

**Table 2** Factor structure

Factor 1 (depression)	Factor 2 (anxiety)	Factor 3 (somatic symptoms)
Crying easily	Felt suddenly scared	Felt weak
Hopeless about future	Felt fearful	No desire for sex
Felt unhappy	Felt nervousness	Poor appetite
Felt without friends	Felt terror or panic	Faint, dizzy or weak
Everything was an effort	Felt difficult to settle	Heart beat so fast
Felt useless		Trembling
Felt tense		Headaches

were classified as having probable depression on the new subscale but not on the Bolton & Ngodoni subscale.

There was a statistically significant inverse correlation between happiness and both the depression ( $r = -0.43$ ; 95% CI  $-0.55$  to  $-0.30$ ) and anxiety ( $r = -0.32$ ; 95% CI  $-0.45$  to  $-0.17$ ) subscales. Cuzick's test confirmed a trend of increasing symptom scores across the different levels of happiness ( $P < 0.001$  for both depression and anxiety). Similarly, there was a statistically significant positive correlation between food insecurity and both the depression ( $r = 0.27$ ; 95% CI  $0.12$ – $0.41$ ) and anxiety ( $r = 0.32$ ; 95% CI  $0.17$ – $0.45$ ) subscales. The somatic symptoms subscale also had a statistically significant inverse correlation with happiness ( $r = -0.21$ ; 95% CI  $-0.35$  to  $-0.05$ ) and a statistically significant positive correlation with food insecurity ( $r = 0.22$ ; 95% CI  $0.07$ – $0.37$ ), although these correlation coefficients were slightly smaller in magnitude. The somatic symptoms subscale had a stronger and statistically significant inverse correlation with self-reported overall health ( $r = -0.40$ ; 95% CI  $-0.53$  to  $-0.26$ ). Cuzick's test confirmed a trend of increasing somatic symptom scores across the different levels of self-reported overall health ( $P < 0.001$ ).

## Discussion

The findings indicate that the HSCL-25 is a reliable and valid measure of depression and anxiety among PLHIV in rural Uganda. Parallel analysis revealed a three-factor solution with three related, but distinct, constructs: depression, anxiety and somatic symptoms. The three subscales were internally consistent. The depression and anxiety sub-scales correlated strongly with happiness and food insecurity while the somatic symptoms sub-scale was more strongly correlated with overall self-reported health, thus providing evidence of construct validity.

The depression and anxiety sub-scales showed slightly higher reliability compared with the somatic symptoms sub-scale. The scale reliability coefficients demonstrated by the depression and anxiety sub-scales are similar in magnitude to those estimated in a previous study of women

recruited from an antenatal clinic in Tanzania [39]. Nearly one-third of participants met criteria for caseness on the depression subscale, which is broadly in agreement with findings from other parts of Uganda and from other countries in sub-Saharan Africa [40–44].

The three-factor structure is similar to what has been found in a related study of PLHIV in Tanzania [17]. The high rate of caseness on the somatic symptoms sub-scale can potentially be explained by the unique features of the sample (i.e., all were HIV positive). Additionally, across different cultures in sub-Saharan Africa, including Uganda, people tend to express their feelings using physical symptoms because they perceive them to be of greater relevance when seeking care [12, 13, 45–47]. Although previous studies from the U.S. have suggested that there is need to exclude somatic symptoms from depression screening tools for PLHIV [16, 48], somatic symptoms are commonly expressed by persons with common mental disorders in most cultures across Uganda. In a previous study of PLHIV in Uganda undergoing treatment for depression, study participants explained depression as an illness of many thoughts and worries that affect physical well-being, potentially explaining the frequent presentation of physical symptoms among them [49]. Therefore, excluding somatic symptoms while assessing for depression among PLHIV may potentially limit the ability to capture symptoms of depression since somatic symptoms have been observed to be one of the principal modes of expressing psychological distress [50, 51].

Our findings should be interpreted in view of several limitations. First, HIV serostatus was based on self-report. Although it is unlikely that many participants were misclassified (i.e., it is unlikely that persons would have reported themselves to be HIV-positive when in fact they were HIV-negative) [20], it is possible that there were some persons in the general study population who were HIV-positive but who were either unaware of their seropositivity or who falsely reported themselves to be HIV-negative (and were therefore not included in this analysis). Both of these are non-trivial possibilities given that many PLHIV in Uganda have not been tested and are unaware of their seropositivity [52] and also given that HIV remains highly

stigmatized in Uganda [53–55]. Nonetheless, our HIV prevalence estimate of 8.7% (based on self-report) is similar to the HIV prevalence estimate of 8.0% for southwest Uganda in the 2011 Uganda AIDS Indicator Survey (based on unlinked anonymous HIV testing) [56]; therefore we anticipate that any potential bias resulting from misclassification would be minimal. Second, we did not have access to a gold standard for the purposes of determining criterion-related validity. Thus, caution should be exercised in reviewing our findings. Studies with population-based data on clinical psychiatric assessments are rare. Future studies validating the HSCL-25 against a gold standard of psychiatric diagnosis among PLHIV should be considered. Third, for the somatic symptoms subscale specifically, we did not have objective measures of disease progression (e.g., CD4 + T-lymphocyte cell count) for our assessments of construct validity. We used a measure of self-reported overall health that is generally accepted as a valid measure of health status [28, 29]. However, because HIV status was self-reported and testing for CD4 levels was not done, we were not able to definitively conclude that the high rate of somatic symptoms was related to advanced HIV disease.

## Conclusions

In summary, this study found that the HSCL-25 is a reliable and valid measure of mental health among PLHIV in rural Uganda. In this population, the prevalence of depression and anxiety are high. Moreover, the experience of depression is a mixed picture of cognitive, affective, anxiety, and somatic symptoms. Excluding somatic symptoms could potentially lead to many undetected cases of depression and, therefore, undertreatment overall. Thus, somatic symptoms should be taken into consideration when assessing for depression among PLHIV in rural Uganda.

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## Compliance with Ethical Standards

**Conflict of interest** We have no conflicts of interest to disclose.

**Ethical Approval** Ethical approval for all study procedures was obtained from the Partners Human Research Committee, Massachusetts General Hospital; and the Institutional Review Committee, Mbarara University of Science and Technology. Consistent with national guidelines, we received clearance for the study from the Uganda National Council for Science and Technology and from the Research Secretariat in the Office of the President.

**Informed Consent** Written informed consent to participate was obtained. Study participants who were unable to write were permitted to indicate consent with a thumbprint.

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