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Structural Validity of the Rheumatology Attitudes Index (RAI) in Systemic Sclerosis: Analysis from the UCLA Scleroderma Quality of Life Study

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Abstract

Objective—To evaluate the structural validity of the Rheumatology Attitudes Index (RAI), a widely used measure of rheumatic disease-related helplessness in patients with systemic sclerosis (SSc).

Methods—Patients with physician-confirmed SSc from the University of California, Los Angeles (UCLA) Scleroderma Quality of Life Study ($N = 208$) received clinical examinations and completed self-report questionnaires. The structural validity of the RAI was examined via confirmatory and exploratory factor analysis (CFA/EFA).

Results—A tenable factor structure was not identified through CFA or EFA.

Conclusion—The present structural analysis did not support the use of the RAI with SSc patients.

Keywords

Helplessness; Systemic sclerosis; Scleroderma; Validity

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In the context of chronic illness, patients may appraise their health as unpredictable and uncontrollable, and believe that their efforts to control their illness will be ineffective (1). In the rheumatic diseases, this cognitive style, termed helplessness, has been associated with greater psychological disability, poorer response to medical treatments, and mortality (2). Helplessness can also mediate the relationship between depression and functional impairment (3). For patients with systemic sclerosis (SSc), a rare rheumatic disease that is characterized by an often unpredictable course of the disease and sudden symptomatic changes (4), helplessness may also be a relevant construct. Previous research examining helplessness in SSc has identified associations between helplessness and depressive symptoms (5, 6).

The 15-item Arthritis Helplessness Index (AHI) was developed to measure helplessness in rheumatoid arthritis (RA) (1). The four-point response options ranged from 1 (*strongly disagree*) to 4 (*strongly agree*). Although the AHI was hypothesized to be unidimensional, in the development study ($N = 219$ patients with RA) the low internal consistency of the total score ($\alpha = 0.69$) suggested a multifactor solution (1). In a follow-up study of RA patients ($N = 368$), explanatory factor analysis (EFA) of the AHI found that five items loaded onto a factor reflecting beliefs that patients *cannot* control disease outcomes, labeled the Helplessness subscale ($\alpha = 0.63$), and six items loaded onto a factor reflecting beliefs that patients *can* control disease outcomes, labeled the Internality subscale ($\alpha = 0.75$) (7). The other items were still retained in the measure; however, they were not included in the scoring. The two subscales were significantly correlated at a small magnitude ($r = 0.21$).

Because the AHI was designed for use in RA, it needed to be adapted to be applied to other rheumatic diseases. The modified version, the Rheumatology Attitudes Index (RAI; 8) is identical to the AHI in content, except for the substitution of “condition” for “arthritis” and a modification of the response scale to include a neutral response option 2.5 (*do not agree or disagree*). Like the AHI, the internal consistency of the RAI total score was marginal ($\alpha = 0.68$); moreover, only the total score was analyzed (8). Additionally, there have been other versions of the RAI response scale, including a six-point response scale (9).

A problem with both the AHI and RAI is their inconsistent application, with some studies using the six-item Internality subscale only, some studies substituting a 7-item Internality subscale, other studies using the Helplessness subscale as a stand-alone measure (10), and still others using the total score of all 15 items (11). Although previous studies have not provided clear rationales for using different variations of the measure, a possible explanation is that the initial RAI development paper (8) was published *prior* to the study demonstrating that the two subscale version of the AHI is psychometrically preferable (7). This timeline may contribute to the inconsistent use of the measure. Another reason may be that, in the same study (7), two different factor solutions were described; a seven-item Internality subscale was found to fit the data in one sample of RA patients ($N = 368$), whereas a six-item Internality subscale was found using a cross-validation sample using the RA patients from the original AHI development study ($N = 219$). Additionally, in a paper describing the measure in a non-RA rheumatic sample, Engle and colleagues (12) stated that the RAI “was initially called the Arthritis Helplessness Index, and was later renamed the Rheumatology Attitudes Index” (p. 281), implying that the latter subsumed the former. Although both

measures have been widely used, many authors using the RAI often refer to the measure as the AHI (e.g., 14). Further details and a summary of the different scoring methods of the measure are available (11).

The RAI has been adopted in several studies in SSc, despite the fact that its structural validity has not been established in this population (6, 13–17). In one of these studies, the five-item Helplessness subscale was used as a standalone measure (6), whereas in the other studies the total score has been used (13–17). Several of these studies (13–16) reference the same paper (12), which described the RAI as a one-factor measure scored by calculating the sum of all 15 items. A formal psychometric evaluation is needed prior to confidently using the RAI in SSc.

METHODS

Patients and Procedure

The sample consisted of 208 patients from the University of California, Los Angeles Scleroderma Quality of Life Study, an observational, single-center, cohort study. Participants were at least 18 years old, and had a formal diagnosis of SSc confirmed by a study physician. The study was approved by UCLA Institutional Review Board (IRB), study number 7-07-061-01.

Measures

Rheumatology Attitudes Index—(RAI; 8). The RAI is a measure of perceived control and helplessness over rheumatic disease-related outcomes. Participants rate a series of 15 statements with responses ranging from 1 (strongly disagree) to 6 (strongly agree). A total score is derived by reverse-scoring nine of the 15 items, and summing the items. Higher scores reflect greater helplessness. Subscale scores for *Helplessness* (5 items) and *Internality* (6 or 7 items) can also be calculated.

Sociodemographic and Medical Characteristics—Patients self-reported sociodemographic details. Disease severity was physician-evaluated using the modified Rodnan skin score (mRSS; 18).

Data Analysis

Descriptive statistics were conducted using SPSS Version 23.0. Confirmatory factor analysis (CFA) was conducted in MPlus version 7.12 to examine the structural validity of AHI/RAI scores. A one-factor structure using all 15 items, both of the two-factor structures (i.e., the five-item Helplessness subscale with either the six-item or seven-item versions of the Internality subscales), and a one-factor Helplessness structure were examined.

Overall model fit was determined using the recommendations of Bentler (19). Three indicators of model fit were utilized: (a) the Root Mean Square Error of Approximation (RMSEA), (b) the Standardized Root Mean Residual (SRMR), and (c) the Comparative Fit Index (CFI). For the RMSEA and SRMR, values less than .08 indicated acceptable fit; for the CFI, values greater than .90 indicated acceptable model fit. A model was determined to

fit well if at least two of these indicators met criteria for acceptable fit. MLR estimation was used in the present analysis. Modification indices were also requested.

In anticipation of the possibility that the CFA would not identify a solution, an EFA was planned. Items with a primary loading $\geq .45$ and secondary loading $\geq .25$ would be retained to maximize practical significance and diminish multi-vocality (20, 21). Parallel analysis was also planned to confirm the number of factors that should be retained in the EFA. The aforementioned descriptive fit indices and theoretical interpretations were considered to make final determinations of factor retention by iteratively removing items.

RESULTS

Descriptive Statistics

Sample characteristics are summarized in Table 1.

Structural Validity

Confirmatory factor analysis—The one-factor total score solution did not fit well statistically ($\chi^2 [53] = 138.80, p < .001$), or descriptively (CFI = .714, RMSEA = .086, SRMR = .077). For the two-factor solution, the Helplessness and Internality latent variables were indicated by five and seven observed items, respectively, as done in the Stein et al. (7) study. This two-factor model did not fit well statistically ($\chi^2 [53] = 138.80, p < .001$), or descriptively (CFI = .706, RMSEA = .086, SRMR = .089). The five-item Helplessness and six-item Internality solution offered by Stein et al. (6) as an alternative was also tested, but did not fit well either statistically ($\chi^2 [43] = 117.63, p < .001$), or descriptively (CFI = .821, RMSEA = .091, SRMR = .068). The one-factor Helplessness scale (5 items) (10) was also tested and did not fit well statistically ($\chi^2 [5] = 12.367, p < .05$), but did demonstrate tenable fit descriptively (CFI = .944, RMSEA = .084, SRMR = .041). However, internal consistency reliability was not adequate ($\alpha = .634$). No modifications were undertaken given a lack of theoretical justification for freeing parameters.

Exploratory factor analysis—Using the .45 factor loading cut-off criterion, the EFA also failed to identify a tenable solution. The model was also evaluated using a less conservative factor loading cut-off of .40. The parallel analysis indicated that a two factor-solution best represented the data when eigenvalues from the present data set were compared to eigenvalues from randomly simulated data: 1) Factor 1: 2.85 vs. 1.33, 2) Factor 2: 1.46 versus 1.22. Using the .40 cut-off, a two-factor solution fit the data per descriptive indices, such that the three descriptive fit indices indicated good fit ($\chi^2 [19] = 22.32, p = .269$; CFI = .985, RMSEA = .029, SRMR = .032). Seven items either did not load at $\geq .40$ onto either factor or loaded at $> .25$ onto both factors and were thus removed, with the exception of one item, described below. Using this approach, nine items were retained (see Table 2), labeled as Helplessness (5 items) and Factor 2 as Internality (4 items). The correlation between the two factors ($r = 0.22$) was similar to a previous study (7). However, this two-factor solution was also deemed untenable because it included an item (item 10) that loaded on both factors, but removing the caused the descriptive fit indices to fall below

acceptable levels. Furthermore, internal consistency reliability was not adequate for either subscale (Helplessness: $\alpha = .634$; Internality: $\alpha = .645$).

DISCUSSION

In the present study of patients with SSc, we did not find support for any of the previously identified factor structures for the RAI through CFA and did not identify an alternative tenable factor structure through EFA. This suggests that the RAI is not an appropriate measure of helplessness in SSc. Brady (11) examined measures of helplessness, self-efficacy, mastery, and control that have been used in rheumatology research. A number of measures of constructs conceptually related to helplessness (e.g., self-efficacy) were identified that may be appropriate for use in SSc, pending psychometric validation.

Limitations of the present study should be considered. The literature contains different response options for the RAI's response scale; the present study used the six-item response scale. Additionally, the sample was limited to patients in Southern California receiving care at a university-hospital setting.

CONCLUSION

The present structural analysis did not support the use of the RAI with SSc patients. In the absence of a valid measure of helplessness for this population, measures of related constructs (e.g., self-efficacy) may be considered.

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Table 1

Demographic and medical variables (N = 208)

	<i>M (SD) or n (%)</i>
Demographic variables	
Age (years)	51.48 (14.28)
Race	
White	149 (71.6%)
Black	13 (6.3%)
Asian	25 (12.0%)
American Indian	5 (2.4%)
Mixed	12 (5.8%)
Missing	4 (2.0%)
Highest level of education	
High School or Less	34 (16.3%)
Some College	73 (35.1%)
College	46 (22.1%)
Graduate Degree	51 (24.5%)
Missing	4 (1.9%)
Annual income	
75,000	77 (37.1%)
> \$75,000.00	71 (34.1%)
Missing/Would rather not say	60 (28.8%)
Sex	
Female	173 (83.2%)
Male	33 (15.9%)
Missing	2 (1.0%)
Relationship Status	
In a relationship	118 (56.7%)
Single/Separated/Divorced	87 (41.9%)
Missing	3 (1.4%)
Disease Subtype	
Limited	105 (50.5%)
Diffuse	83 (39.9%)
Overlap	7 (3.4%)
Missing or Unknown	13 (6.2%)
Medical variables	
Years since onset of first non-Raynaud's symptom	8.45 (6.73)
Years since diagnosis of SSc	7.57 (7.89)
mRSS score	8.66 (8.51) [max = 51]

Table 2

Factor loadings of the Helplessness and Internality Factors from the Exploratory Factor Analysis Nine-Item Solution

Item (original RAI item number) ^a	Helplessness Factor	Internality Factor
1. My condition is controlling my life. (1)	.592	.048
2. I can reduce my pain by staying calm and relaxed. (3)	.051	.424*
3. If I do all the right things, I can successfully manage my condition. (5)	.133	.431*
4. When I manage my personal life well, my condition does not flare up as much. (8)	-.004	.712*
5. I have considerable ability to control my pain. (9)	.187	.534*
6. I would feel helpless if I couldn't rely on other people for help with my condition. (10)	.717*	-.427^a
7. No matter what I do or how hard I try, I just can't seem to get relief from my pain. (12)	.637*	.033
8. I am coping effectively with my condition. (13)	.462	.102
9. It seems that fate and other factors beyond my control affect my condition. (15)	.457	-.056

Note.

* $p < 0.05$.

^a Dropped items included the following: Item 2: Managing my condition is largely my own responsibility; Item 4: Too often my pain just seems to hit me out of the blue; Item 6: I can do a lot of things myself to cope with my condition; Item 7: When it comes to condition, I feel I can only do what my doctor tells me to do; Item 11: Usually, I can tell when my condition is going to flare up.