

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

Shadi Gholizadeh, Sarah D. Mills, Rina S. Fox, Erin L. Merz, Scott C. Roesch, Philip J. Clements, Suzanne Kafaja, Daniel E. Furst, Dinesh Khanna, and Vanessa L. Malcarne

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 through 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. (J Rheumatol First Release April 15 2017; doi:10.3899/jrheum.161080)

Key Indexing Terms:

HELPLESSNESS SYSTEMIC SCLEROSIS ASSESSMENT INDICES VALIDITY

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 ineffective¹. In the rheumatic diseases, this cognitive style, called “helplessness,” has been associated with greater psychological

disability, poorer response to medical treatments, and mortality². Helplessness can also mediate the relationship between depression and functional impairment³. For patients with systemic sclerosis (SSc), a rare rheumatic disease that is characterized by an often unpredictable disease course and sudden symptomatic changes⁴, helplessness may also be a relevant construct. Previous research examining helplessness in SSc has identified associations between helplessness and depressive symptoms^{5,6}.

From the San Diego State University (SDSU)/University of California (UC) San Diego Joint Doctoral Program in Clinical Psychology, Department of Psychology, San Diego; Department of Psychology, California State University, Dominguez Hills; SDSU, Department of Psychology, San Diego; David Geffen School of Medicine, University of California, Los Angeles (UCLA) School of Medicine, Los Angeles, California; University of Michigan Health System, Ann Arbor, Michigan, USA.

Supported by the grant, Evaluation of Health-Related Quality of Life in Systemic Sclerosis from the Scleroderma Foundation Inc. Dr. Khanna has been funded by the US National Institutes of Health (NIH)/US National Institute of Arthritis and Musculoskeletal and Skin Diseases grants K24 AR063120 and K23 AR053858. Dr. Khanna has served as a consultant for Bayer and Genentech.

S. Gholizadeh, Doctoral Student, MS, MSc, MPH, SDSU/UC San Diego Joint Doctoral Program in Clinical Psychology, Department of Psychology; S.D. Mills, Doctoral Student, MS, MPH, SDSU/UC San Diego Joint Doctoral Program in Clinical Psychology, Department of Psychology; R.S. Fox, Postdoctoral Fellow, PhD, MPH, SDSU/UC San Diego Joint Doctoral Program in Clinical Psychology, Department of Psychology; E.L. Merz, Assistant Professor, PhD, MPH, California State University, Dominguez Hills, Department of Psychology; S.C. Roesch, Professor, PhD, SDSU, Department of Psychology; P.J. Clements, MD, Professor Emeritus, David Geffen School of Medicine, UCLA; S. Kafaja, MD, Clinical Instructor, David Geffen School of Medicine, UCLA; D.E. Furst, MD, Professor, David Geffen School of Medicine, UCLA; D. Khanna, MD, Professor, University of Michigan Health System; V.L. Malcarne, PhD, Professor, SDSU/UC San Diego Joint Doctoral Program in Clinical Psychology, and Department of Psychology, SDSU. Address correspondence to Professor V.L. Malcarne, Doctoral Training Facility, 6363 Alvarado Court, Suite 103, San Diego, California 92120-4913, USA. E-mail: vmalcarne@mail.sdsu.edu

Accepted for publication February 7, 2017.

The 15-item Arthritis Helplessness Index (AHI) was developed to measure helplessness in rheumatoid arthritis (RA)¹. The 4-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 multifactorial solution¹. In a followup study of patients with RA (n = 368), explanatory factor analysis (EFA) of the AHI found 5 items loaded onto a factor reflecting beliefs that patients cannot control disease outcomes, labeled the Helplessness subscale ($\alpha = 0.63$), and 6 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 but they were not included in the scoring. The 2 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 analyzed⁸. Additionally, there have been other versions of the RAI response scale, including a 6-point response scale⁹.

A problem with both the AHI and RAI is their inconsistent application, with some studies using the 6-item Internality subscale only, some studies substituting a 7-item Internality subscale, other studies using the Helplessness subscale as a standalone measure¹⁰, and still others using the total score of all 15 items¹¹. Although previous studies have not provided clear rationales for using different variations of the measure, a possible explanation is that the initial RAI development paper⁸ was published prior to the study, demonstrating that the 2-subscale version of the AHI is psychometrically preferable⁷. This timeline may contribute to the inconsistent use of the measure. Another reason may be that, in the same study⁷, 2 different factor solutions were described: a 7-item Internality subscale was found to fit the data in 1 sample of patients with RA ($n = 368$), whereas a 6-item Internality subscale was found using a cross-validation sample with the patients with RA from the original AHI development study ($n = 219$). Additionally, in a paper describing the measure in a non-RA rheumatic sample, Engle and colleagues¹² stated that the RAI “was initially called the Arthritis Helplessness Index, and was later renamed the Rheumatology Attitudes Index,” 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., McNearney, *et al*¹³). Further details and a summary of the different scoring methods of the measure are available¹¹.

The RAI has been adopted in several studies in SSc, even though its structural validity has not been established in this population^{6,13,14,15,16,17}. In one of these studies, the 5-item Helplessness subscale was used as a standalone measure⁶, whereas in the other studies the total score has been used^{13,14,15,16,17}. Several of these studies^{13,14,15,16} reference the same paper¹², which described the RAI as a 1-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.

MATERIALS AND METHODS

Patients and procedure. The sample consisted of 208 patients from the University of California, Los Angeles (UCLA) 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 the UCLA Institutional Review Board, study number 7-07-061-01.

Measures. 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 9 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 score¹⁸.

Data analysis. Descriptive statistics were analyzed 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 1-factor structure using all 15 items, both of the 2-factor structures (i.e., the 5-item Helplessness subscale with either the 6-item or 7-item versions of the Internality subscales), and a 1-factor Helplessness structure were examined.

Overall model fit was determined using the recommendations of Bentler¹⁹. Three indicators of model fit were used: (1) the root mean square error of approximation (RMSEA), (2) the standardized root mean residual (SRMR), and (c) the comparative fit index (CFI). For the RMSEA and SRMR, values < 0.08 indicated acceptable fit; for the CFI, values > 0.90 indicated acceptable model fit. A model was determined to fit well if at least 2 of these indicators met criteria for acceptable fit. Maximum likelihood estimation with robust standard errors (i.e., 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 ≥ 0.45 and secondary loading ≤ 0.25 would be retained to maximize practical significance and diminish multivocality^{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. In conducting confirmatory factor analysis, the 1-factor total score solution did not fit well statistically ($\chi^2 [53] = 138.80$, $p < 0.001$) or descriptively (CFI = 0.714, RMSEA = 0.086, SRMR = 0.077). For the 2-factor solution, the Helplessness and Internality latent variables were indicated by 5 and 7 observed items, respectively, as in the Stein, *et al*⁷ study. This 2-factor model did not fit well statistically ($\chi^2 [53] = 138.80$, $p < 0.001$), or descriptively (CFI = 0.706, RMSEA = 0.086, SRMR = 0.089). The 5-item Helplessness and 6-item Internality solution offered by Stein, *et al*⁷ as an alternative was also tested, but did not fit well statistically ($\chi^2 [43] = 117.63$, $p < 0.001$) or descriptively (CFI = 0.821, RMSEA = 0.091, SRMR = 0.068). The 1-factor Helplessness scale (5 items)¹⁰ was also tested and did not fit well statistically ($\chi^2 [5] = 12.367$, $p < 0.05$), but did demonstrate tenable fit descriptively (CFI = 0.944, RMSEA = 0.084, SRMR = 0.041). However, internal consistency reliability was not adequate ($\alpha = 0.634$). No modifications were undertaken given a lack of theoretical justification for freeing measures.

Exploratory factor analysis. Using the 0.45-factor-loading cutoff criterion, the EFA also failed to identify a tenable solution. The model was also evaluated using a less-conservative factor loading cutoff of 0.40. The parallel analysis indicated that a 2-factor-solution best represented the data when eigenvalues from the present dataset were compared to eigenvalues from randomly simulated data: (1) Factor 1: 2.85 versus 1.33, (2) Factor 2: 1.46 versus 1.22. Using the ≤ 0.40

Table 1. Demographic and medical variables (n = 208). Data are n (%) unless otherwise indicated.

Demographic Variables	Values
Age, yrs, mean (SD)	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)
Education	
High school or less	34 (16.3)
Some college	73 (35.1)
College graduate	46 (22.1)
Graduate degree	51 (24.5)
Missing	4 (1.9)
Annual income (US\$)	
≤ \$75,000	77 (37.1)
> \$75,000	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, mean (SD)	
Years since first non-Raynaud symptom	8.45 (6.73)
Years since diagnosis of SSc	7.57 (7.89)
mRSS score	8.66 (8.51; max = 51)

SSc: systemic sclerosis; mRSS: modified Rodnan skin score.

cutoff, a 2-factor solution fit the data per descriptive indices, such that the 3 descriptive fit indices indicated good fit (χ^2 [19] = 22.32, p = 0.269; CFI = 0.985, RMSEA = 0.029, SRMR = 0.032). Seven items either did not load at ≥ 0.40 onto either factor or loaded at > 0.25 onto both factors and were thus removed, with the exception of 1 item, described below. Using this approach, 9 items were retained (Table 2), labeled as Helplessness (5 items) and factor 2 as Internality (4 items). The correlation between the 2 factors (r = 0.22) was similar to that in a previous study⁷. However, this 2-factor solution was also deemed untenable because it included an item (item 10) that loaded on both factors, but removing it caused the descriptive fit indices to fall below acceptable levels. Further, internal consistency reliability was not adequate for either subscale (Helplessness: α = 0.634; Internality: α = 0.645).

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. Brady¹¹ 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 6-item response scale. Additionally, the sample was limited to patients in Southern California receiving care at a university-hospital setting.

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.

Table 2. Factor loadings of the Helplessness and Internality factors from the exploratory factor analysis 9-item solution.

Item (original RAI item number) ^a	Helplessness Factor	Internality Factor
1. My condition is controlling my life. (1)	0.592	0.048
2. I can reduce my pain by staying calm and relaxed. (3)	0.051	0.424
3. If I do all the right things, I can successfully manage my condition. (5)	0.133	0.431
4. When I manage my personal life well, my condition does not flare up as much. (8)	-0.004	0.712
5. I have considerable ability to control my pain. (9)	0.187	0.534
6. I would feel helpless if I couldn't rely on other people for help with my condition. (10)	0.717	-0.427
7. No matter what I do or how hard I try, I just can't seem to get relief from my pain. (12)	0.637	0.033
8. I am coping effectively with my condition. (13)	0.462	0.102
9. It seems that fate and other factors beyond my control affect my condition. (14)	0.457	-0.056

^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 my 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; Item 15: I want to learn as much as I can about my condition.

REFERENCES

1. Nicassio PM, Wallston KA, Callahan LF, Herbert M, Pincus T. The measurement of helplessness in rheumatoid arthritis: the development of the Arthritis Helplessness Index. *J Rheumatol* 1985;12:462-7.
2. Jensen MP, Turner JA, Romano JM, Karoly P. Coping with chronic pain: a critical review of the literature. *Pain* 1991;47:249-83.
3. Nicassio PM, Schuman C, Radojevic V, Weisman MH. Helplessness as a mediator of health status in fibromyalgia. *Cognit Ther Res* 1999;23:181-96.
4. Thombs BD, van Lankveld W, Bassel M, Baron M, Buzza R, Haslam S, et al. Psychological health and well-being in systemic sclerosis: state of the science and consensus research agenda. *Arthritis Care Res* 2010;62:1181-9.
5. Kwakkenbos L, van Lankveld WG, Vonk MC, Becker ES, van den Hoogen FH, van den Ende CH. Disease-related and psychosocial factors associated with depressive symptoms in patients with systemic sclerosis, including fear of progression and appearance self-esteem. *J Psychosom Res* 2012;72:199-204.
6. Matsuura E, Ohta A, Kanegae F, Haruda Y, Ushiyama O, Koarada S, et al. Frequency and analysis of factors closely associated with the development of depressive symptoms in patients with scleroderma. *J Rheumatol* 2003;30:1782-7.
7. Stein MJ, Wallston KA, Nicassio PM. Factor structure of the Arthritis Helplessness Index. *J Rheumatol* 1988;15:427-32.
8. Callahan LF, Brooks RH, Pincus T. Further analysis of learned helplessness in rheumatoid arthritis using a "Rheumatology Attitudes Index". *J Rheumatol* 1988;15:418-26.
9. Nicassio PM, Ormseth SR, Custodio MK, Olmstead R, Weisman MH, Irwin MR. Confirmatory factor analysis of the Pittsburgh Sleep Quality Index in rheumatoid arthritis patients. *Behav Sleep Med* 2014;12:1-12.
10. Devellis RF, Callahan LF. A brief measure of helplessness in rheumatoid disease: the helplessness subscale of the Rheumatology Attitudes Index. *J Rheumatol* 1993;20:866-6.
11. Brady TJ. Measures of self-efficacy, helplessness, mastery, and control: The Arthritis Helplessness Index (AHI)/Rheumatology Attitudes Index (RAI), Arthritis Self-Efficacy Scale (ASES), Children's Arthritis Self-Efficacy Scale (CASE), Generalized Self-Efficacy Scale (GSES), Mastery Scale, Multi-Dimensional Health Locus of Control Scale (MHLC), Parent's Arthritis Self-Efficacy Scale (PASE), Rheumatoid Arthritis Self-Efficacy Scale (RASE), and Self-Efficacy Scale (SES). *Arthritis Care Res* 2003;49:S147-64.
12. Engle EW, Callahan LF, Pincus T, Hochberg MC. Learned helplessness in systemic lupus erythematosus: analysis using the Rheumatology Attitudes Index. *Arthritis Rheum* 1990;33:281-6.
13. McNearney TA, Reveille JD, Fischbach M, Friedman AW, Lisse JR, Goel N, et al. Pulmonary involvement in systemic sclerosis: associations with genetic, serologic, sociodemographic, and behavioral factors. *Arthritis Care Res* 2007;15:318-26.
14. McNearney TA, Hunnicutt SE, Fischbach M, Friedman AW, Aguilar M, Ahn CW, et al. Perceived functioning has ethnic-specific associations in systemic sclerosis: another dimension of personalized medicine. *J Rheumatol* 2009;36:2724-32.
15. Reveille JD, Fischbach M, McNearney T, Friedman AW, Aguilar MB, Lisse J, et al. Systemic sclerosis in 3 US ethnic groups: a comparison of clinical, sociodemographic, serologic, and immunogenetic determinants. *Semin Arthritis Rheum* 2001; 30:332-46.
16. Sharif R, Mayes MD, Nicassio PM, Gonzalez EB, Draeger H, McNearney TA, et al. Determinants of work disability in patients with systemic sclerosis: a longitudinal study of the GENISOS cohort. *Semin Arthritis Rheum* 2011;41:38-47.
17. Callahan LF, Smith WJ, Pincus T. Self-report questionnaires in five rheumatic diseases comparisons of health status constructs and associations with formal education level. *Arthritis Rheum* 1989;2:122-31.
18. Clements PJ, Lachenbruch PA, Ng SC, Simmons M, Sterz M, Furst DE. Skin score: a semiquantitative measure of cutaneous involvement that improves prediction of prognosis in systemic sclerosis. *Arthritis Rheum* 1990;33:1256-63.
19. Bentler PM. On tests and indices for evaluating structural models. *Pers Individ Dif* 2007;42:825-9.
20. Clark LA, Watson D. Constructing validity: basic issues in objective scale development. *Psychol Assess* 1995;7:309-19.
21. Tabachnick BG, Fidell LS, Osterlind SJ. Using multivariate statistics, 4th ed. New York: Harper & Row; 2001.