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Psychometric analysis and validation of the Persian translation of the systemic sclerosis questionnaire (SySQ)

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Abstract:

BACKGROUND: Scleroderma is a complex multisystem disorder that could have effects on the quality of patients' lives. This study was conducted by determining the psychometric properties of the Persian version of the systemic sclerosis questionnaire (SySQ) that specifically assesses indications and functional limitations of scleroderma patients.

MATERIALS AND METHODS: In the present cross-sectional study, the method included: translation and back translation. Psychometric properties of the questionnaire including its content and face validity were assessed. Internal consistency with the SySQ (Cronbach's alpha) and reproducibility was by test-retest method. The factor structure of the questionnaire was evaluated using exploratory factor analysis. The convergent validity of the SySQ was assessed using the General Health Assessment Scale (HAQ).

RESULTS: Altogether 32 SySQ items, the internal consistency coefficient (Cronbach's alpha) of the whole tool was 0.906. The content validity index was 0.98 and the content validity ratio was 0.796, there was a significant relationship between the questions and the relevant factors in the factor analysis. The correlation coefficient = 0.953 for the instrument. The correlation of SySQ dimensions with HAQ questionnaire dimensions in convergent validity showed that musculoskeletal dimensions, general condition, and cardiorespiratory of the questionnaire are correlated with all dimensions of the HAQ questionnaire.

CONCLUSION: The Persian version of the questionnaire SySQ with competency is valid and reliable and is suitable for measuring specific changes in Persian systemic sclerosis patients.

Keywords:

Psychometrics, reliability, systemic sclerosis, validity

Introduction

The quality of life of scleroderma patients can be affected by the disease process. Scleroderma^[1] is a complex multisystem disorder.^[2] It is a chronic disease with an unknown cause and autoimmune origin, which is associated with excessive collagen secretion and connective tissue problems^[3] and is characterized by fibroblast dysfunction. This disease affects the walls of blood vessels, as well as the skin of the internal organs of the body.^[4,5]

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It is crucial to note that scleroderma is a long-term disease^[6] that progressively affects various organs of the body.^[7] Consequently, it leads to severe disabilities in the physical and mental social functioning of patients, such as the inability to work and participate in family life, the disruption of personal hygiene, the fear of disease progression, and dissatisfaction with the body image because skin stiffness leads to changes in the appearance,^[1] digital wounds, and oral and dental problems. In addition, the involvement of the gastrointestinal system, shortness of breath, general pain and

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0.618 after two weeks.^[34] The similarity of the results obtained in these studies can be due to the specificity of the questionnaire in terms of the symptoms and challenges of scleroderma patients.

In this study, the questionnaire has been evaluated, with wider and more comprehensive dimensions, (content validity, face validity, construct validity, convergent validity, and reliability) compared to other studies, and more importantly, the results have been from good to excellent, and this provides the strength and validity of the Persian questionnaire. But other translations have examined the questionnaire in a more limited way and in someone have adopted relatively lower results.

So, the Persian version of this questionnaire is eligible for evaluation and shows the condition, specific symptoms, and functional limitations of scleroderma patients. This questionnaire can be effective in reducing the complications of the disease, maintaining and improving the patient's recovery at any level of the disease. Therefore, the author hopes that the findings of this research will help to a better quality of life, development of management, care, and educational and targeted treatment programs in various fields including nursing of disease.

Limitations and recommendation

One of the limitations of this disease is that it is in the category of rare diseases with few facilities and support, and due to poor health, weakness, and weakness of the immune system in Corona conditions, it makes it difficult for patients to participate in research. In addition, it is challenging to obtain samples due to the limited number of treatment centers dedicated to these patients. In addition, this questionnaire does not specifically assess important factors such as Raynaud's phenomenon and kidney problems.

Conclusion

The aim of this study was to conduct a psychometric analysis of the Persian version of the SYSQ effective on quality of life questionnaire in scleroderma patients. The findings of the study showed that this questionnaire has good content validity, construct validity, and reliability and can be useful in the assessment of Persian-speaking patients. Using this tool, the challenges and the process of the disease and its treatment can be examined more accurately. The findings of this study can be helpful to researchers, nursing managers, and nurses. This study can be used as a basis for further studies to pave the way for the evaluation of these patients by considering other characteristics of the disease and adding wider domains of the patient's issues, including kidney problems and Raynaud's phenomenon, to the questionnaire.

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Conflicts of interest

There are no conflicts of interest.

References

1. Alian SM, Sherby NA, Sarhan SA. Cultural adaptation and validation of the systemic sclerosis quality of life questionnaire into Arabic language. *Clin Rheumatol* 2021;40:1409-16.
2. Barsotti S, Orlandi M, Codullo V, Di Battista M, Lepri G, Della Rossa A, *et al.* One year in review 2019: Systemic sclerosis. *Clin Exp Rheumatol* 2019;37(Suppl 119):3-14.
3. Lachner KD. Caring for the patient with limited systemic sclerosis. *Orthop Nurs* 2016;35:5-10.
4. Hinkle JL, Cheever KH. Brunner and Suddarth's textbook of medical-surgical nursing: Wolters Kluwer India Pvt Ltd; 2018.
5. Machado RI, Souto LM, Freire EA. Translation, cultural adaptation and validation into Portuguese (Brazil) in Systemic Sclerosis Questionnaire (SySQ). *Rev Bras Reumatol* 2014;54:95-101.
6. Kocher A, Ndosi M, Denhaerynck K, Simon M, Dwyer AA, Distler O, *et al.* A rare disease patient-reported outcome measure: Revision and validation of the German version of the systemic sclerosis quality of life questionnaire (SScQoL) using the Rasch model. *Orphanet J Rare Dis* 2021;16:356.
7. Sandqvist G, Hesselstrand R. Validity of the Swedish version of the systemic sclerosis quality of life questionnaire (SScQoL): A novel measure of quality of life for patients with systemic sclerosis. *Ann Rheum Dis* 2019;78:855-7.
8. Merz EL, Malcarne VL, Roesch SC, Nair DK, Salazar G, Assassi S, *et al.* Longitudinal patterns of pain in patients with diffuse and limited systemic sclerosis: Integrating medical, psychological, and social characteristics. *Qual Life Res* 2017;26:85-94.
9. Pauling JD, Reilly E, Smith T, Frech TM. Factors influencing Raynaud condition score diary outcomes in systemic sclerosis. *J Rheumatol* 2019;46:1326-34.
10. Pope JE, Bellamy N. Outcome measurement in scleroderma clinical trials. *Semin Arthritis Rheum* 1993;23:22-33.
11. Tugwell P, Boers M, Brooks P, Simon L, Strand V, Idzerda L. OMERACT: An international initiative to improve outcome measurement in rheumatology. *Trials* 2007;8:38.
12. Sierakowska M, Doroszkiewicz H, Sierakowska J, Olesińska M, Grabowska-Jodkowska A, Brzosko M, *et al.* Factors associated with quality of life in systemic sclerosis: A cross-sectional study. *Qual Life Res* 2019;28:3347-54.
13. Fries JF, Spitz PW, Young DY. The dimensions of health outcomes: The health assessment questionnaire, disability and pain scales. *J Rheumatol* 1982;9:789-93.
14. Poole JL, Steen VD. The use of the health assessment questionnaire (HAQ) to determine physical disability in systemic sclerosis. *Arthritis Care Res* 1991;4:27-31.