RELIABILITY AND STABILITY OF THE ROLAND MORRIS DISABILITY QUESTIONNAIRE:
INTRA CLASS CORRELATION AND LIMITS OF AGREEMENT

CHAPTER 2

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Chapter 2

ABSTRACT

Objective: To analyse test-retest reliability and stability of the Dutch language version of the Roland Morris Disability Questionnaire (RMDQ-Dv) in a sample of patients (n=30) suffering from Chronic Low Back Pain (CLBP).

Design: Test-retest reliability was studied after the patients filled out the RMDQ-Dv twice with a two-week interval and before starting the rehabilitation program. Intra Class Correlation (ICC), (one way random) was used as a measure for reliability and the limits of agreement were calculated for quantifying the stability. An ICC of 0.75 or more was considered as an acceptable reliability. No criteria for limits of agreement were available. However, smaller limits of agreement indicate more stability because it indicates that the natural variation is small.

Subjects: Thirty patients (24 male and 6 female) with CLBP participated in this study. All patients were referred for treatment in a rehabilitation centre between May 2000 and April 2001 and agreed to participate. Demographics and medical history were obtained of all patients. The mean age of the patients was 40 years (SD 8.1). The duration of low back pain ranged between 5 and 10 years. Patients were off work for a mean of 17 weeks (SD 19.2). Fifteen patients (50%) were receiving financial compensation.

Results: The RMDQ-Dv showed good reliability, with an ICC of 0.91. Calculating limits of agreement to quantify the stability, a large amount of natural variation (± 5.4) was found relative to the total scoring range of 0 to 24.

Conclusion: The RMDQ-Dv proves to be a reliable instrument to measure functional status in CLBP patients. However, the natural variation should be taken into account when using it clinically.
Reliability and stability of the RMDQ

INTRODUCTION

Functional status is an important evaluative outcome measure in low back pain rehabilitation.\textsuperscript{1,2} To assess changes in functional status after treatment in patients with low back pain, the Roland Morris Disability Questionnaire (RMDQ) is frequently used.\textsuperscript{2-4} The RMDQ is derived from the Sickness Impact Profile, a general health questionnaire.\textsuperscript{5}

To assess functional status, it is important that the reliability of the instrument is good and that repeated measures in individuals remain stable over time,\textsuperscript{6} in the absence of treatment. In reliability studies of the RMDQ, Pearson correlation coefficient is often used as a measure for reliability.\textsuperscript{2,7,8} Pearson correlation reflects the extent to which two repeated measures can be fitted by a straight line. The disadvantage of this statistic measure is that repeated measures may differ systematically (statistically), yet correlate highly or perfectly. By contrast, the intra-class correlation coefficient (ICC) assesses not only the strength of correlation, but also if all measures on each subject are identical, and do not differ systematically. Therefore, ICC is preferable over the Pearson correlation to use as measure for reliability. But usually the Pearson coefficient will be higher than the ICC and may be used more often for that reason.

Stability over time, in the absence of treatment, may be influenced by within-patient variance and random errors. These sources of variance may lead to instability or fluctuations on the RMDQ-scale: ‘natural variations’.\textsuperscript{6} If a person fills out the same questionnaire on two occasions, it is relevant to know what variation in test scores can be expected in the absence of treatment. To investigate this natural variation on the RMDQ-scale, limits of agreement can be calculated according to the method of Bland and Altman.\textsuperscript{9} In an individual patient the change due to treatment should exceed these limits of agreement before one can state that the treatment has been effective. Therefore, limits of agreement should be taken into account when using the RMDQ clinically.

The English language version of the RMDQ shows good reliability.\textsuperscript{1,7,10} However, limits of agreement have not been investigated. A validated Dutch language version of the RMDQ (RMDQ-Dv) is available,\textsuperscript{11} but test-retest reliability and limits of agreement have not been investigated previously.

The aim of this study is to investigate the test-retest reliability of the RMDQ-Dv in patients with chronic low back pain (CLBP), using ICC as measure for reliability, as well as to quantify the stability of the RMDQ-Dv by calculating limits of agreement.
METHODS

General procedure
Patients with CLBP were recruited from the population who were admitted for rehabilitation treatment of the Centre for Rehabilitation at the University Hospital Groningen. Patients were included in the study if they were between 18-65 years of age, still at work, or were less than 1 year out of work due to CLBP. Exclusion criteria were specific low back pain, entirely off work for a year or more, cardiovascular or pulmonary diseases, pregnancy, addiction, and psychopathology. Patients filled out the RMDQ-Dv twice, before starting the rehabilitation program, with a two-week interval. Time, day and place of assessment were held constant for the two test-sessions. The present study was approved by the Medical Ethical Committee of the University Hospital Groningen.

Subjects
Thirty patients (24 male and 6 female) with CLBP participated in this study. All patients were referred for treatment in a rehabilitation centre between May 2000 and April 2001 and agreed to participate. Demographics and medical history were obtained of all patients. The mean age of the patients was 40 years (SD 8.1). The duration of low back pain ranged between 5 and 10 years. Patients were off work for a mean of 17 weeks (SD 19.2). Fifteen patients (50%) were receiving financial compensation.

RMDQ-Dv
The RMDQ-Dv is a translation of the original RMDQ.\textsuperscript{8} It assesses perceived limitations in 24 activities of daily living dichotomously. The sum score is calculated by summing the "yes" answers. The scale ranges from 0 (no disability) to 24 (severe disability).

Data analyses
Descriptive statistics were calculated for the total scores of the two test-sessions. Test-retest reliability was determined by means of a paired t-test, Intra Class Correlation (ICC, one way random) for the sum scores. Limits of agreement were used to determine the natural variation for quantifying stability over time.\textsuperscript{9,12} To calculate limits of agreement, a plot of the difference between the two sessions for each patient against the mean of each patient of the two sessions was made. Then the average difference in the two sessions, and the standard deviation of the difference between the two scores (SDchange) were calculated. Finally, the limits of agreement were calculated, equal to twice the standard deviation. An ICC above 0.75 was considered as good reliability.\textsuperscript{13,14} No criteria for interpretation of the limits of agreement were available. However, smaller limits of agreement indicate more stability because it indicates that the natural variation is small. Data
analyses were performed using the Statistical Package for Social Sciences (SPSS 10.0).

RESULTS

Mean of the sum score in the first and second session was respectively 13.0 (SD 4.8) and 12.1 (SD 5.0). The mean difference was 0.83 (SD 2.7) (95% CI of the difference: -0.2 to 1.8). The ICC was 0.91 (95% CI: 0.82 to 0.96). Limits of agreement were ± 5.4 (figure 1).

Figure 1. Difference between RMDQ1 and RMDQ 2 plotted against average of RMDQ1 and RMDQ2
DISCUSSION

No systematic differences were found in the sum score of the first and the second session. The reliability of the RMDQ-Dv was good (ICC (one way random) above the criterion of 0.75). Similar results of ICCs of 0.75 or higher were found in many other RMDQ studies.10,16-21 However, also considerable lower ICCs were found ranging from 0.42 to 0.66.21,22 Most studies with ICC values of ≥ 0.75 used an interval of 1-14 days between the two sessions, whereas for the studies with ICCs below 0.75, the interval was 6 weeks or more. Almost all studies with a time interval of more than two weeks have lower ICCs than the studies with an interval of two weeks or less. An explanation of this phenomenon might be that a shorter interval between the two sessions may result in patients remembering the score of the previous session. A larger interval between the two sessions may result in loss of remembering the score of the previous session and change of clinical status in that period. Reliability of functional status questionnaires may be best measured using an interval of 1-2 weeks, a period in which the clinical status is reasonably stable in chronic pain patients.4 In our study we used an interval of two weeks.

Comparing studies using Pearson correlation2,7,8 with studies using ICC10,16-21 as a measure of reliability, it appears that the magnitude of Pearson and ICC are similar, i.e. the reliability is good. This suggests that the predominant source of error is due to random variation instead of a systematic difference. Under these circumstances, the Pearson and ICC are very similar.14

To quantify stability, we investigated the natural variation by calculating limits of agreement according to the method of Bland and Altman.9 Despite the good reliability (ICC), the limits of agreement (± 5.4) were large relative to the total scoring range of 0 to 24. This means that within-patient variance or random errors have led to instability in measurement results, approximately 95% of all differences within persons will lie between ± 5.4. This large amount of natural variation should be taken into account when using the RMDQ-Dv clinically. Effects of therapy should exceed the limits of agreement before one can state that the treatment has been effective. Post-hoc analysis showed that for all items ≥ 70% of the scores were the same for the two sessions. Thus, the large amount of natural variation could not be contributed to some specific items.

De Vet et al.6 used the Smallest Real Differences for individuals (SRDindividual) as measure for quantifying the stability of the RMDQ-Dv. Despite the use of different terms, the calculation of the limits of agreement and SRDindividual are the same. We found limits of agreement of 5.4 on a scale of 0-24, de Vet et al.6 found a SRDindividual value of 5.9. Limits of agreement, in our study, were calculated on the basis of scores collected before patients started the intervention, to minimize the possibility that a clinically important change of the construct
would occur in the period of data collecting. The study of de Vet et al.,\(^6\) however, is an intervention study and the SRD\(_{\text{individual}}\) was calculated on the basis of the scores of a group of patients who rated themselves as not clinically important changed despite the intervention. An estimation of not clinically important changed was obtained by global perceived effect assessed by the patient on a 7-points transition scale (1 = completely recovered, 7 = vastly worsened).\(^{23}\) The validity of a transition scale however is problematic; it cannot be regarded as a 'golden standard'. Bias may affect subjective assessment of change, patients do not remember correctly how they felt at the beginning of treatment. Moreover, they usually underestimate their initial state, resulting in exaggerating the effect of the program.\(^{14}\) Classifying patients as "changed" or "not-changed" on the basis of these results may be biased.

Limits of agreement can also be used as a cut-off score for change in an intervention study or in daily practice. The cut-off change score determines the minimum change that is considered to be clinically relevant.\(^{24,25}\) Based on our results, patients have to change at least 6 points on a scale of 0-24 of the RMDQ-Dv to exceed the natural variation and to be judged as having really changed. Several intervention studies have determined the cut-off score for change of the RMDQ,\(^ {4,21,26,27}\) ranging from 2 to 5 points. These studies underestimate the height of the cut-off score; changes on the RMDQ scale ranging from 2 to 5 points cannot be detected as a clinically relevant change, given the natural variation we found.

**CONCLUSION**

The RMDQ-Dv proves to be a reliable instrument to assess functional status in CLBP patients. However, a large amount of natural variation (± 5.4) was found relative to the total scoring range of 0 to 24.
REFERENCES


