

Title: Test and Re-test of the painDETECT Questionnaire

R. Baron^a, R. Freynhagen^c, U. Gockel^d, T. Kohlmann^e, T. Keller^b, E. Stemmler^f and T.R. Tölle^g

^a Sektion Neurologische Schmerzforschung und Therapie, Universitätsklinikum Schleswig-Holstein, Campus Kiel, Germany

^b StatConsult GmbH, Magdeburg, Germany

^c Zentrum für Anästhesiologie, Intensivmedizin, Schmerztherapie & Palliativmedizin, Benedictus-Krankenhaus, Tutzing, Germany

^d Grünenthal GmbH, Aachen, Germany

^e Institute for Community Medicine, Universität Greifswald, Germany

^f Pfizer Pharma GmbH, Berlin, Germany

^g Technische Universität München, Klinik für Neurologie, Munich, Germany

Background and Objective

Since its introduction in 2006 the painDETECT questionnaire (PD-Q) has been widely used. Data from more than 225,000 patients have been collected in the painDETECT project register. The original validation did not include 'test-retest' because of the resulting necessity to suspend or interrupt pain treatment.

Validation of the test-retest performance of PD-Q items and the derived PD-Q score is reported.

Methods

For the patients in the data base with stable disease, retrospective analysis of 2 consecutive visits was performed. To ensure stable disease, the visits had to fulfil the following criteria: interval between visits, 7–21 days; time since first capture in database, ≥ 6 months; indication, back pain; difference of average, greatest and current pain between visits each less than 5 points on the 100-point NRS.

It was verified that the selected sub-population was representative of the whole study population.

Intra-class-correlation ICC, Pearson correlation and weighted kappa were used as statistical measures for ordinal scaled items. Bland–Altman plot and Passing–Bablok regression were also assessed for continuously scaled PD-Q score.

Results

Data from 94 patients fulfilled the very narrow criteria; mean duration between visits was 15 days. There was no relevant deviation of mean PD-Q score or pain severity in comparison with the whole study population. The measures were in the range of typical results for pain questionnaires (e.g. VAS).

	PD-Q Items	PD-Q score
ICC	0.65 – 0.80	0.87
Pearson's r	0.66 – 0.80	0.87
Weighted kappa	0.50 – 0.66	–
BA plot	–	Mean(SD) of difference: –0.36 (3.85)
PB regression	–	Slope = 1 Intercept = 0

Conclusions

This validation has shown that the PD-Q is reliable and can be used for follow-up. Further investigations will be necessary to determine the clinical relevance of changes in PD-Q score.