Reliability and Validity of the Korean Short-form Eight-item Parkinson’s Disease Questionnaire (PDQ-8)

Sehyun Kim1, Hye-Ja Park2, Min Jung Park3, Hyun Sook Kim4

1Professor, Graduate School, Dankook University, Jukjeon; 2Professor, College of Nursing, CHA University, Pocheon; 3Professor, Department of Nursing, CHA Bundang Medical Center, CHA University, Seongnam; 4Professor, Department of Neurology, CHA Bundang Medical Center, CHA University, Seongnam, Korea

**Objectives:** This study aimed to determine the reliability and validity of the Korean Parkinson’s disease questionnaire (PDQ-8) using a standard short-form validation approach. **Methods:** This cross-sectional psychometric study evaluated 80 outpatients with PD who completed the Korean 39-item Parkinson’s disease questionnaire (PDQ-39) and PDQ-8 tools in random order. The summary index (SI) was calculated for the PDQ-39 (PDQ-39SI) and PDQ-8 (PDQ-8SI). A neurologist performed evaluations using the Hoehn and Yahr (HY) scale, unified Parkinson’s disease rating scale (UPRDS motor), mini-mental state examination (MMSE), Montreal cognitive assessment (MoCA), the clinical dementia rating (CDR), and global deterioration scale (GDS). **Results:** The PDQ-8SI exhibited appropriate floor effect (2.5%) and ceiling effect (1.3%), with acceptable internal consistency reliability (Cronbach’s α = 0.88), item-total correlations (0.48-0.77), and inter-rater reliability (Kappa coefficient = 0.53). The PDQ-8 loaded one factor (Eigen value = 4.35, 54.51% of the variance) and had broadly similar factor loadings (0.585-0.847). The PDQ-8 had mostly goodness of fit (GFI = 0.932, NFI = 0.914, TLI = 0.973, IFI = 0.981, CFI = 0.981, and RMSEA = 0.056). The PDQ-8SI and PDQ-8 items exhibited good correlations with each other (all p < 0.05) and the PDQ-39SI (r = 0.91, p < 0.001). The PDQ-8SI was correlated with levodopa dose (r = 0.28, p = 0.028), HY score (r = 0.44, p = 0.001), UPRDS motor (r = 0.45, p = 0.007), MMES (r = 0.28, p = 0.047), MoCA (r = -0.49, p = 0.006), GDS (r = 0.33, p = 0.025), and CDR (r = 0.40, p = 0.007), respectively. **Conclusions:** The PDQ-8 appears to be reliable and valid. It may be useful for quickly and easily evaluating the patient’s perspectives regarding the disease’s impacts, which may guide nursing practice and future research.

**Key words:** Parkinson’s disease, Quality of life, Reliability, Validity

INTRODUCTION

Parkinson’s disease (PD) is one of the three major geriatric neurodegenerative diseases and is an important issue among Koreans [1]. During the last 10 years, this disease has exhibited annual increases of 13.2% in its incidence and 21% in related healthcare costs [1,2]. Patients with PD primarily experience motor symptoms, including resting tremors, rigidity, and bradykinesia, which are caused by pathological loss or degeneration of dopaminergic neurons in the substantia nigra of the midbrain, as well as the development of neuronal Lewy bodies [3]. As the disease progresses, patients also develop non-motor symptoms, including cognitive, mood, and behavioral difficulties, which significantly increase their overall disability and significantly decrease their health-related quality of life (HRQoL) [3,4].

Clinicians have access to conventional instruments for assessing PD severity and disability, such as the Hoehn and Yahr scale (HY scale) [5] and the unified Parkinson’s disease rating scale [6]. However, these assessments do not provide information regarding the overall health and disease’s impacts from the patient’s perspective [7,8]. Nevertheless, HRQoL is based on the patient-reported outcomes of the physical, psy-
chological, and social effects of their disease [9,10]. Importance of HRQoL assessment has increased to provide life-extending treatment and supportive care for PD patients and their families [11]. Thus, reliable and valid HRQoL measures are essential for understanding the multifaceted effects of PD and are crucial for promoting patient-centered integrated healthcare throughout the course of PD [12,13].

The 39-item Parkinson’s disease questionnaire (PDQ-39) has been widely translated and validated in multiple languages as a core PD-specific HRQoL instrument [7,8,10,14]. This instrument evaluates the effects of PD on HRQoL based on the patient’s perspectives covering mobility, activities of daily living, emotional well-being, stigma, social support, cognition, communication, and bodily discomfort [14]. However, the PDQ-39 takes some time to complete leading to the administration burden on respondents and its use has been restricted primarily to research settings or less advanced patients [8]. Therefore, to reduce respondent burden, a shorter 8-item PDQ (PDQ-8) has been constructed by selecting the most highly correlated item from each dimension of the PDQ-39 [8,14]. As a brief and simple measure, the PDQ-8 is more suitable than the PDQ-39 for patients to quickly and easily complete in waiting rooms before they see their doctors or health care providers [7,8]. Implementing such a practice may improve health care provider-patient communications, patients’ satisfaction, medical decision making, and eventually, patients’ wellbeing [7,8]. Summary score can prove helpful in providing a guide on the overall impact of ill health as measures on questionnaires [7,14]. The PDQ-8 provides a summary index score of impact of PD on self-reported quality of life among PD patients [7,14]. Moreover, the PDQ-8 have been reflected the health condition including Hoehn and Yahr stage of the disease severity as a gold standard [5,7,8,14], the motor function [6,8], the cognitive ability [15,16], and dementia [17,18]. Thus, the PDQ-8 permits more economical screening the patient’s health status, and it has been increasingly used in clinical trials and in everyday healthcare settings [7,14].

Previous studies validated the PDQ-8 using the PDQ-8 items “nested” within the existing PDQ-39 data [7,14,19], despite the standard validation practice of the short form instrument should be performed by administering the instrument as an independent instrument [8,19]. This validation approach may influence patient responses and its psychometric properties [8], and a standard validation with the short-form measure is needed to determine whether the PDQ-8 is a useful independent instrument [7,8,19]. However, limited studies have been performed using a standard short-form validation approach with the PDQ-8 [7,8,19]. Furthermore, the Korean PDQ-8 has not been validated using a standard short-form validation approach. The present study aimed to evaluate the reliability and validity of the Korean PDQ-8 as an independent instrument for use in clinical nursing practice and research.

METHODS

Setting and participants

This cross-sectional study was conducted at a university hospital-affiliated neurology outpatient center, where PD patients visited from throughout the nation. The G*Power 3.1.7 program was used to estimate the optimal sample size, which was determined to be 26 patients for determining convergent validity according to the point-biserial correlation model (a two-sided test, $\alpha = 0.05$, power of $1-\beta = 0.80$, and effect size $\rho = 0.50$) based on the previous studies [8,20]. To use factor analysis for determining the instrument’s psychometric properties, we multiplied the 8 items (PDQ-8) or 8 scale-based dimensions (PDQ-39) by 10 to reach a number of 80. Eighty patients with neurologist-diagnosed PD were enrolled from July 2014 to June 2016. All participant were cognitively able to read and complete the questionnaires. Patients with a history of neurological diseases other than PD and psychiatric disorders were excluded based on the neurologist’s decision. We administered the Korean PDQ-39 and PDQ-8 instruments to the 80 consecutive outpatients after they had provided informed consent. Thirty six (45.0%) consecutively filled out the PDQ-8 and PDQ-39, and 44 (55.0%) consecutively completed the PDQ-39 and PDQ-8 questionnaires in random order. To minimize missing data, a clinical research nurse checked for completeness when each patient returned his or her completed questionnaire at a counseling room. All the participants were included in this study without any dropouts. The validation procedures were based on the guidelines of the original authors and the international validation steps [9,14] with the 8-dimension of the PDQ-39, the nested PDQ-8 extracted from the PDQ-39 and the independently administered PDQ-8 tools. A neurologist measured disease severity, motor function, cognitive function, and degree of dementia and then recorded the results in the electronic medical record system. This study protocol and procedures were approved by the institutional review board at Bundang CHA Medical Center (IRB No. BD2014-104).
Measurements

Demographics and clinical assessment

The participants’ demographic characteristics were collected using a self-administered questionnaire regarding age, employment status, and marital status. We also reviewed their electronic medical records to obtain data regarding clinical characteristics.

The HY scale estimates disease severity (0-5) and is the gold standard for assessing clinical outcomes [5]. The Unified Parkinson Disease Rating Scale (UPRDS) motor scale measures motor function (0-33) with higher scores indicating poorer motor function [6]. The Mini-Mental State Examination (MMSE) simply assesses the patient’s cognitive function (0-30) with higher scores indicating better cognitive function [15]. The Montreal Cognitive Assessment (MoCA) sensitively screens for cognitive deficits (0-30), with higher scores indicating better cognitive function [16]. The Geriatric Depression Scale (GDS) measures the stage of dementia (1-5) with higher scores indicating poorer dementia condition [17]. The Clinical Dementia Rating scale (CDR) also measures the stage of dementia, with higher scores indicating poorer dementia ranging from CDR 0 (no dementia) to CDR 5 (terminal dementia) [18].

The Parkinson’s disease questionnaires (PDQ-39 and PDQ-8), and PDQ summary index

The original author approved the use of the British English PDQ-39 and PDQ-8 and provided a user manual [14]. The use of the Korean PDQ-39 was also approved by the author [20]. The Korean PDQ-39 was developed through the steps of translation (forward translation and backward translation), cognitive debriefing by a group of specialists, a pilot test, and a transcultural adaptation test [21]. The Korean PDQ-39 exhibited the adequate content, construct, and criterion validity and reliability. Cronbach’s α of the 8 dimension ranged from 0.70 to 0.97 in the previous study [20]. Data from the PDQ-39 or PDQ-8 can be presented in profile form as well as summarized in the PDQ-39 summary index (PDQ-39SI) or PDQ-8 summary index (PDQ-8SI) [14].

The PDQ-39 instrument consists of eight dimensions: mobility, activities of daily living, emotional well-being, stigma, social support, cognition, communication, and bodily discomfort. All items are scored from 0 (never) to 4 (always), and each dimension is transformed using a scale from 0 (no problem) to 100 (maximum difficulty). The PDQ-39 summary index (PDQ-39SI) is calculated by summing the 8-dimension scores and then standardizing the score on a scale of 0–100, with lower scores indicating better HRQoL. [14] (Table 1).

The PDQ-8 includes eight specific items selected based on the strongest correlations for the eight dimensions of the PDQ-39. The eight items are scored from 0 (never) to 4 (always), and then a summary index of the PDQ-8 (PDQ-8SI) is calculated based on the eight scores using a standardized scale (0–100), with lower scores indicating better HRQoL. [14] (Table 1).

Data analysis

All analyses were performed using IBM SPSS 23.0 software (IBM Corp., Armonk, NY, USA) and AMOS 24 software (IBM Corp., Armonk, NY, USA).

(1) Data normality was determined using the Kolmogorov-Smirnov test.

(2) Differences in the mean scores between the PDQ-39SI, the nested PDQ-8SI, and the independent PDQ-8SI were analyzed using the Kruskal-Wallis test with Bonferroni’s method for multiple comparisons.

(3) Floor and ceiling effects were examined based on the response ranges and distributions for each item.

(4) Internal consistency reliability was assessed using Cronbach’s α and the item-total correlations. The inter-rater reliability was evaluated using the Kappa coefficient between the nested PDQ-8 items and the independent PDQ-8 items.

(5) Exploratory and confirmatory factor analyses were used to evaluate construct validity.

(6) For convergent validity, the Spearman’s correlation analysis was performed to evaluate the relationship among the independent PDQ-8 items, as well as with the PDQ-8SI and PDQ-39SI.

(7) For the criterion validity, the Spearman’s correlation analysis was used to evaluate the relationship among the independent PDQ-8SI, levodopa dose, HY stage, UPRDS III, MMSE, MoCA, GDS, and CDR scores.

RESULTS

Demographics and clinical assessment

The mean age was 67.55 ± 9.31 years and 51 patients (63.7%) had an onset age of ≥ 65 years. Female accounted for 49 (61.2%). The mean disease duration was 14.18 ± 28.65 months. The mean levodopa dose was
Floor and ceiling effects, and reliability

The PDQ-8SI exhibited the appropriate response ranges and distributions for each item, with a 2.5% floor effect and a 1.3% ceiling effect. The internal consistency reliability of the independent PDQ-8 was good.
The item-total correlations ranged from 0.48 to 0.77, with 0.53 inter-rater reliability (Kappa coefficient) between the independent and nested PDQ-8 versions ($p < 0.001$). Cronbach’s $\alpha$ of the PDQ-39 was 0.97. Cronbach’s $\alpha$ of the 8 dimensions in the PDQ-39 ranged from 0.72 to 0.94 (Table 3).

**Content validity**

The conceptual equivalence and consistency between the British English PDQ-8 and Korean PDQ-8 instrument was evaluated by two neurology nursing specialists, a nursing professor, and a neurologist. The content validity index (CVI) was acceptable (CVI = 0.95).

**Construct validity**

The means for the independent PDQ-8 items were generally similar to the means for the 8 dimensions in the PDQ-39 and the nested PDQ-8, except for the “activities of daily living” item ($p = 0.047$) (Table 4). The independent PDQ-8 was loaded on one factor (Eigen value = 4.35, 54.51% of the variance), with broadly similar factor loadings (0.585–0.847). The independent PDQ-8 had mostly goodness of fit (GFI = 0.932, 0.908).

---

**Table 2. Demographics and clinical assessment (n=80)**

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>Categories</th>
<th>Mean ± SD or n (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sociodemographic</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Age (y)</td>
<td></td>
<td>67.55 ± 9.31</td>
</tr>
<tr>
<td>Onset age</td>
<td>&lt; 65</td>
<td>29 (36.3)</td>
</tr>
<tr>
<td></td>
<td>≥ 65</td>
<td>51 (63.7)</td>
</tr>
<tr>
<td>Sex</td>
<td>Male</td>
<td>31 (38.8)</td>
</tr>
<tr>
<td></td>
<td>Female</td>
<td>49 (61.2)</td>
</tr>
<tr>
<td>Employment status</td>
<td>Employed</td>
<td>12 (15.0)</td>
</tr>
<tr>
<td></td>
<td>Unemployed</td>
<td>68 (85.0)</td>
</tr>
<tr>
<td>Spouse</td>
<td>Yes</td>
<td>51 (63.7)</td>
</tr>
<tr>
<td></td>
<td>No</td>
<td>29 (36.3)</td>
</tr>
</tbody>
</table>

Clinical indices
- Duration of disease (mon) 14.18 ± 28.65
- Levodopa dose (mg) (n = 63) 370.24 ± 258.25
- Hoehn & Yahr staging score (n = 51) 1.56 ± 1.14
- UPDRS motor (n = 34) 25.74 ± 17.10
- MMES (n = 50) 25.54 ± 3.16
- MoCA (n = 31) 18.00 ± 8.22
- GDS (n = 46) 2.67 ± 1.17
- CDR (n = 44) 0.93 ± 0.96

SD, standard deviation; UPDRS, unified Parkinson’s disease rating scale; MMES, mini-mental state examination; MoCA, Montreal cognitive assessment; GDS, global deterioration scale; CDR, clinical dementia rating; PDQ, Parkinson’s disease questionnaire.

**Table 3. Means, floor effect, ceiling effect, and reliability (n=80)**

<table>
<thead>
<tr>
<th>Dimensions (Item number of PDQ-8)</th>
<th>Mean ± SD</th>
<th>Independent PDQ-8</th>
<th>$\chi^2$</th>
<th>$p$</th>
<th>Cronbach’s $\alpha$</th>
<th>Item-total-correlation</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>PDQ-39</td>
<td>Nested PDQ-8</td>
<td>PDQ-8</td>
<td></td>
<td>PDQ-39</td>
<td>Nested PDQ-8</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PDQ-39</td>
<td>46.53 ± 30.19</td>
<td>46.25 ± 36.26</td>
<td>41.56 ± 34.21</td>
<td>1.16</td>
<td>0.560</td>
<td>0.94</td>
</tr>
<tr>
<td>Activities of daily living (Q2)</td>
<td>39.22 ± 30.86</td>
<td>32.19 ± 37.14</td>
<td>35.63 ± 36.84</td>
<td>6.10</td>
<td>0.047</td>
<td>0.92</td>
</tr>
<tr>
<td>Emotional well-being (Q3)</td>
<td>41.61 ± 25.77</td>
<td>42.82 ± 29.55</td>
<td>40.00 ± 28.57</td>
<td>0.44</td>
<td>0.802</td>
<td>0.93</td>
</tr>
<tr>
<td>Social support (Q4)</td>
<td>21.25 ± 22.26</td>
<td>24.06 ± 29.08</td>
<td>24.38 ± 32.80</td>
<td>0.94</td>
<td>0.626</td>
<td>0.81</td>
</tr>
<tr>
<td>Cognitions (Q5)</td>
<td>40.73 ± 24.77</td>
<td>39.06 ± 32.03</td>
<td>36.88 ± 33.51</td>
<td>1.35</td>
<td>0.509</td>
<td>0.72</td>
</tr>
<tr>
<td>Communication (Q6)</td>
<td>23.44 ± 24.42</td>
<td>23.13 ± 27.78</td>
<td>23.13 ± 29.97</td>
<td>1.37</td>
<td>0.505</td>
<td>0.82</td>
</tr>
<tr>
<td>Bodily pain (Q7)</td>
<td>33.02 ± 25.45</td>
<td>38.44 ± 30.28</td>
<td>40.94 ± 30.87</td>
<td>2.93</td>
<td>0.231</td>
<td>0.77</td>
</tr>
<tr>
<td>Stigma (Q8)</td>
<td>25.52 ± 24.96</td>
<td>27.50 ± 28.01</td>
<td>27.81 ± 30.55</td>
<td>0.05</td>
<td>0.976</td>
<td>0.84</td>
</tr>
<tr>
<td>Summary index (SI)</td>
<td>33.92 ± 24.40</td>
<td>34.18 ± 22.44</td>
<td>33.79 ± 23.65</td>
<td>0.19</td>
<td>0.908</td>
<td></td>
</tr>
<tr>
<td>% Floor</td>
<td>1.3</td>
<td>2.5</td>
<td>2.5</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>% Ceiling</td>
<td>1.3</td>
<td>1.3</td>
<td>1.3</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Kappa coefficient</td>
<td></td>
<td>0.53 ($p &lt; 0.001$)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>


Kappa coefficient between the nested PDQ-8 items and the independent PDQ-8 items.

Bonferroni multiple comparisons, the same letter are not statistically significant.
Table 4. Construct validity based on the exploratory and confirmatory factor analyses (n=80)

<table>
<thead>
<tr>
<th>Dimensions</th>
<th>Factor loadings by EFA</th>
<th>β (p) by CFA</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>PDQ-39</td>
<td>Nested PDQ-8</td>
</tr>
<tr>
<td>Mobility</td>
<td>0.813</td>
<td>0.707</td>
</tr>
<tr>
<td>Activities of daily living</td>
<td>0.810</td>
<td>0.809</td>
</tr>
<tr>
<td>Emotional well-being</td>
<td>0.780</td>
<td>0.687</td>
</tr>
<tr>
<td>Social support</td>
<td>0.827</td>
<td>0.849</td>
</tr>
<tr>
<td>Cognitions</td>
<td>0.686</td>
<td>0.653</td>
</tr>
<tr>
<td>Communication</td>
<td>0.827</td>
<td>0.674</td>
</tr>
<tr>
<td>Bodily pain</td>
<td>0.768</td>
<td>0.579</td>
</tr>
<tr>
<td>Stigma</td>
<td>0.735</td>
<td>0.765</td>
</tr>
</tbody>
</table>

Keiser-Meyer-Olkin 0.87 0.84 0.81
Bartlett’s χ² (p) 378.15 (0.001) 265.18 (0.001) 277.52 (0.001)
Eigen value 4.89 4.15 4.35
% of variance explained 61.17 51.86 54.51

Absolute fit indices
χ² (CMIN) 32.02 34.05 25.03
df 18 19 20
p 0.022 0.018 0.200
Q (CMIN/df) 1.779 1.792 1.252
GFI 0.915 0.907 0.932
RMR 0.042 0.088 0.074

Relative fit indices
NFI 0.924 0.877 0.914
TLI 0.945 0.911 0.973
CFI 0.965 0.942 0.981
RMSEA 0.964 0.940 0.981

Noncentrality indices
CFI 0.964 0.940 0.981
RMSEA 0.999 0.999 0.056

PDQ, Parkinson’s disease questionnaire; EFA, exploratory factor analysis; CFA, confirmatory factor analysis; CMIN, minimum chi-square; df, degree of freedom; Q, normed chi-square; GFI, goodness of fit index; NFI, normed fit index; RMR, root mean square residual; TLI, Tucker-Lewis index, non-normed fit index; IFI, incremental fit index; CFI, comparative fit index; RMSEA, root mean square error of approximation.

NFI = 0.914, TLI = 0.973, IFI = 0.981, CFI = 0.981, and RMSEA = 0.056 (Table 4).

Convergent validity and criterion validity

For convergent validity, the eight items of the independent PDQ-8 were correlated with each other (all p < 0.05), as well as between the independent PDQ-8SI and PDQ-39SI scores (r = 0.91, p < 0.001) (Table 5).

For criterion validity, the independent PDQ-8SI was correlated with the levodopa dose (r = 0.28, p = 0.028) and the HY score (r = 0.44, p = 0.001). The independent PDQ-8SI was correlated with the scores for UPDRS motor (r = 0.45, p = 0.007), MMSE (r = -0.28, p = 0.047), MoCA (r = -0.49, p = 0.006), GDS (r = 0.33, p = 0.025), and CDR (r = 0.40, p = 0.007) (Table 5).

DISCUSSION

Assessment of health-related quality of life (HRQOL) is essential in developing and promoting best practices for Parkinson’s disease (PD) care [8,10]. The 8-item Parkinson’s disease questionnaire (PDQ-8) provides a global measure of health and is accurately representative of the parent 39-item PDQ (PDQ-39) [7,14,19]. Two methods were used for the linguistic validation and cross-cultural adaptation of the PDQ-8 [8,14,20-23]. The first one is to use the 8 items derived from the existing PDQ-39 data [7,14,19]. The other is to standard validation practice to independently administer the short-form questionnaire to respondents [8,21-23].

In this study, the floor and ceiling effects were both judged appropri-
Table 5. Spearman correlation coefficients for convergent validity and criterion validity (n=80)

<table>
<thead>
<tr>
<th>Independent PDQ-8 (n=80)</th>
<th>Q1</th>
<th>Q2</th>
<th>Q3</th>
<th>Q4</th>
<th>Q5</th>
<th>Q6</th>
<th>Q7</th>
<th>PDQ-39SI</th>
<th>Nested PDQ-8SI</th>
<th>Independent PDQ-8SI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Q1</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>0.62 (&lt;0.001)</td>
<td>0.66 (&lt;0.001)</td>
<td>0.73 (&lt;0.001)</td>
</tr>
<tr>
<td>Q2</td>
<td>0.51 (&lt;0.001)</td>
<td>0.56 (&lt;0.001)</td>
<td>0.57 (&lt;0.001)</td>
<td>0.61 (&lt;0.001)</td>
<td>0.65 (&lt;0.001)</td>
<td>0.66 (&lt;0.001)</td>
<td>0.63 (&lt;0.001)</td>
<td>0.69 (&lt;0.001)</td>
<td>0.74 (&lt;0.001)</td>
<td></td>
</tr>
<tr>
<td>Q3</td>
<td>0.39 (&lt;0.001)</td>
<td>0.31 (0.005)</td>
<td>0.37 (0.001)</td>
<td>0.55 (&lt;0.001)</td>
<td>0.57 (&lt;0.001)</td>
<td>0.60 (&lt;0.001)</td>
<td>0.68 (&lt;0.001)</td>
<td>0.70 (&lt;0.001)</td>
<td>0.76 (&lt;0.001)</td>
<td></td>
</tr>
<tr>
<td>Q4</td>
<td>0.62 (&lt;0.001)</td>
<td>0.57 (&lt;0.001)</td>
<td>0.37 (0.001)</td>
<td>0.68 (&lt;0.001)</td>
<td>0.70 (&lt;0.001)</td>
<td>0.76 (&lt;0.001)</td>
<td>0.61 (&lt;0.001)</td>
<td>0.62 (&lt;0.001)</td>
<td>0.69 (&lt;0.001)</td>
<td></td>
</tr>
<tr>
<td>Q5</td>
<td>0.31 (0.005)</td>
<td>0.54 (&lt;0.001)</td>
<td>0.32 (0.004)</td>
<td>0.50 (&lt;0.001)</td>
<td>0.53 (&lt;0.001)</td>
<td>0.68 (&lt;0.001)</td>
<td>0.70 (&lt;0.001)</td>
<td>0.70 (&lt;0.001)</td>
<td>0.70 (&lt;0.001)</td>
<td></td>
</tr>
<tr>
<td>Q6</td>
<td>0.47 (&lt;0.001)</td>
<td>0.63 (&lt;0.001)</td>
<td>0.32 (0.004)</td>
<td>0.53 (0.001)</td>
<td>0.53 (&lt;0.001)</td>
<td>0.68 (&lt;0.001)</td>
<td>0.61 (&lt;0.001)</td>
<td>0.62 (&lt;0.001)</td>
<td>0.69 (&lt;0.001)</td>
<td></td>
</tr>
<tr>
<td>Q7</td>
<td>0.34 (0.002)</td>
<td>0.41 (&lt;0.001)</td>
<td>0.25 (0.024)</td>
<td>0.32 (0.004)</td>
<td>0.28 (0.012)</td>
<td>0.33 (0.003)</td>
<td>0.58 (&lt;0.001)</td>
<td>0.57 (&lt;0.001)</td>
<td>0.58 (&lt;0.001)</td>
<td></td>
</tr>
<tr>
<td>Q8</td>
<td>0.43 (&lt;0.001)</td>
<td>0.51 (&lt;0.001)</td>
<td>0.47 (&lt;0.001)</td>
<td>0.65 (&lt;0.001)</td>
<td>0.48 (&lt;0.001)</td>
<td>0.47 (&lt;0.001)</td>
<td>0.39 (&lt;0.001)</td>
<td>0.67 (&lt;0.001)</td>
<td>0.70 (&lt;0.001)</td>
<td>0.74 (&lt;0.001)</td>
</tr>
</tbody>
</table>

PDQ-8SI

| PDQ-8SI | 0.91 (<0.001) | 0.94 (<0.001) | 0.91 (<0.001) |

Clinical indices

| Levodopa dose (n = 63) | 0.32 (0.011) | 0.32 (0.011) | 0.28 (0.028) |
| Hoehn and Yahr score (n = 51) | 0.38 (0.005) | 0.44 (0.001) | 0.44 (0.001) |
| UPDRS motor (n = 34) | 0.50 (0.003) | 0.51 (0.002) | 0.45 (0.007) |
| MMES (n = 50) | -0.33 (0.018) | -0.32 (0.024) | -0.28 (0.047) |
| MoCA (n = 31) | -0.50 (0.004) | -0.49 (0.005) | -0.49 (0.006) |
| GDS (n = 46) | 0.37 (0.010) | 0.38 (0.010) | 0.33 (0.025) |
| CDR (n = 44) | 0.49 (0.001) | 0.46 (0.002) | 0.40 (0.007) |

PDQ, Parkinson's disease questionnaire; SI, summary index; HY, Hoehn & Yahr; UPDRS, unified Parkinson's disease rating scale; MMES, mini-mental state examination; MoCA, Montreal cognitive assessment; GDS, global deterioration scale; CDR, clinical dementia rating.

ate based on an acceptable setting points of < 20% [24], as well as findings from previous studies [8,20-23]. The internal consistency reliability should be evaluated for measuring the strength of agreement between repeated measures [9]. In this study, the independent PDQ-8 revealed acceptable internal consistency reliability (α = 0.88) and the item-total correlations (0.48-0.73) based on the acceptable points of Cronbach’s α of ≥ 0.7 and the item-total correlation coefficient of ≥ 0.20 [9]. This result may be supported by the consistent previous reports [8,14,19,21-23]. The Kappa coefficient demonstrated a moderate inter-rater reliability between the independent and nested PDQ-8 instruments. This result contrasts with the outstanding Kappa coefficient of 0.983 from a previous study of 114 PD patients [22]. The discrepancy may indicate a disagreement in patient responses between the nested and independent PDQ-8 findings (the lower activities of daily living (ADL) score in the nested PDQ-8 relative to the independent PDQ-8) [7,8,19]. However, the Korean PDQ-8 is a reliable and reproducible independent instrument.

The data quality for HRQoL depends on the completeness of scale-level and item-level data [10]. The present study completely collected the PDQ-39 and the PDQ-8 items, and the mean scores for the PDQ-8SI and PDQ-39SI were similar to the scores from a multi-national study involving America, Canada, Japan, Italy, Spain [19], and Korea [20]. The independent PDQ-8 had an one-dimensional construct with acceptable factor loadings (≥ 0.40) [9], as well as previous reports [8,19,22]. In the confirmatory factor analysis, the independent PDQ-8 had no discrepancy between the sample and fitted covariance matrices based on the χ² result (p > 0.05) and the model Q (GFI > 0.20) for a good fit. The independent PDQ-8 had a GFI of 0.932, which suggests the appropriate proportion of variance accounted for by the estimated population covariance. However, the RMR value was barely appropriate (0.074) based on recommended cut-offs of < 0.05 for the best fit point and < 0.08 for a good fit point. In contrast, the RMR of the nested PDQ-8 did not meet the good fit point (0.088), despite the RMR of the PDQ-39 being considered excellent (0.042). The relative fit indices (NFI, TLI, IFI) were acceptable based on the conventional cut point of ≥ 0.9, CFI ≥ 0.9, and RMSEA < 0.08, which means all measured variables are uncorrelated. In this context, the Korean PDQ-8 has a one-dimensional construct and appears to be valid.

The eight items of the Korean PDQ-8 were correlated with each other as well as with the PDQ-39SI, the nested PDQ-8SI, and the independent PDQ-8SI based on a minimum acceptable convergent validity value of 0.4 [9]. This result indicates that the independent Korean PDQ-8 has good convergent validity.

The Korean PDQ-8SI was correlated with HY staging score and UPRDS motor score as clinical gold standards, although all the participants did not complete those measurements [8,14,19]. Even mild cognitive impairment associated with Parkinson’s disease (PD) is a risk factor for the development of dementia and HRQoL deterioration, which highlights the importance of early identification of mild cognitive impairment in PD [3,11,12]. The independent PDQ-8SI was also correlated with cognitive function based on the MMSE [15] and MoCA scores [16], and dementia stage based on the GDS [17] and CDR [18]. Therefore, the independent Korean PDQ-8 appears to have good criterion validity. These findings may help nurses assess overall health based on physical, emotional, cognition, and social aspects. Furthermore, the assessment is relatively rapid (5–10 min) and may facilitate the best possible care.

The present study has several limitations that warrant consideration. First, although the study was performed with a sample of PD patients who visited a single university hospital from throughout the nation, the findings may not necessarily generalize to other regions or populations. Second, the sample size (n = 80) is considered barely adequate for comparing the psychometric properties between the PDQ-39 and the PDQ-8, despite the use of scale-level or item-parceling factor analysis for the PDQ-39. Third, the present study enrolled patients with relatively mild disease severity, which is associated with a risk of bias, therefore future studies are needed to test this issue in a larger and more diverse group of patients with PD.

**CONCLUSION**

The results of the present study suggest that the independent Korean PDQ-8 may be a reliable and valid instrument for evaluating HRQoL among PD patients. This tool may be useful for simply and rapidly assessing the effects of PD on patients’ lives, based on their perspective rather than a clinical assessment, which may help guide everyday nursing practice and PD-related research.

**ORCID**

Kim Sehyun  https://orcid.org/0000-0002-1931-7169

Hye-Ja Park  https://orcid.org/0000-0002-8923-2611
REFERENCES


국문초록
한국어판 파킨슨병 삶의 질 축약형 도구의 신뢰도와 타당도
김세현1, 박혜자2, 박민정3, 김현숙4
1단국대학교 대학원 교수, 2차의과학대학교 간호대학 교수, 3차의과학대학교 분당차병원 간호국 교수, 4차의과학대학교 분당차병원 신경과 교수

목적: 파킨슨병 환자를 위한 건강관리 전략을 개발하고 증진하는데 건강 관련 삶의 질 측정은 매우 중요하다. 8개의 항목으로 구성된 파킨슨병 삶의 질 도구(PDQ-8)는 39개 항목으로 개발된 파킨슨병 삶의 질 도구(PDQ-39)의 축약형으로 파킨슨병 환자의 건강상태를 측정하는데 유용하다. 본 연구는 한국어판 축약형 파킨슨병 삶의 질 측정도구의 신뢰도와 타당도를 확인하고자 한다.


결과: 축약형 파킨슨병 삶의 질 도구의 천장효과 2.5%와 바닥효과 1.3%는 적절하였고 내적일관성 신뢰도 계수(α=0.88), 문항-총점 상관 계수(0.48-0.77), 캐스틱관계계수 0.53은 적절하였다. 축약형 파킨슨병 삶의 질 문항은 하나의 요인에 적합되었으며(고윳값=4.35, 분산율=54.51%), 요인적재량은 만족되었으며(0.585-0.847). 확인적요인분석의 모형은 적합하였다(GFI=0.932, NFI=0.914, TLI=0.971, IFI=0.981, CFI=0.981, RMSEA=0.056). 축약형 파킨슨병 삶의 질 도구는 문항 간 유의한 상관관계를 보였고(p<0.05). PDQ-8 건강상태지수는 PDQ-39 건강상태지수와 유의한 상관관계를 보였고(r=0.91, p<0.001) 수렴타당도는 만족되었다. 축약형 파킨슨병 삶의 질 건강상태 지수는 레보도파민 복용량(r=0.28, p=0.028), 파킨슨병 중증도(r=0.44, p=0.001), 운동능력(r=0.45, p=0.007), 간이정신검사(r=0.44, p=0.047), 몬트리올 인지평가(r=0.49, p=0.006), 임상지매척도(r=0.33, p=0.025), 국제치매척도(r=0.40, p=0.007)와 유의한 상관관계를 보여 준거타당도를 만족하였다.

결론: 한국어판 축약형 파킨슨병 삶의 질 도구는 파킨슨병 환자가 가각하는 건강상태를 측정할 수 있는 신뢰도와 타당도가 높은 도구로 간호실무와 연구 수행에 유용한 도구로 사용할 수 있을 것이다.

주제어: 파킨슨병, 삶의 질, 신뢰도, 타당도