Validation of a UPDRS-/MDS-UPDRS-based definition of functional dependency for Parkinson’s disease

Neil Ramsay MRCPa1, Angus D Macleod MRCP PhDa2, Guido Alves MD PhDb,c, Marta Camacho MSd, Lars Forsgren MD PhDe, Rachael A Lawson PhDf, Jodi Maple-Grødem PhDg,b,c Ole-Bjørn Tysnes MD PhDg, Caroline H Williams-Gray MRCP PhDd, Alison J Yarnall MRCP PhDf, Carl E Counsell MRCP MDg on behalf of the Parkinson’s Incidence Cohorts Collaboration

a. Institute of Applied Health Sciences, University of Aberdeen, Aberdeen, UK
b. The Norwegian Centre for Movement Disorders, Stavanger University Hospital, Norway
c. Department of Chemistry, Bioscience and Environmental Engineering, University of Stavanger, Norway
d. Department of Clinical Neurosciences, University of Cambridge, UK
e. Department of Clinical Science, Neurosciences, Umeå University, Sweden
f. Translational and Clinical Research Institute, Newcastle University, UK
g. Department of Neurology, Haukeland University Hospital, University of Bergen, Norway

1Joint first authors
*Corresponding Author: Room 1.015, Polwarth Building, Foresterhill Campus, University of Aberdeen, AB25 2ZD, Email: angus.macleod@abdn.ac.uk Phone: +44 (0)1224 437120

Email addresses of authors:
Dr Ramsay: neil.ramsay@nhs.net
Dr Macleod: angus.macleod@abdn.ac.uk
Prof Alves: guided.werner.alves@sus.no
Marta Camacho: msc72@cam.ac.uk
Prof Forsgren: Lars.Forsgren@umu.se
Dr Lawson: rachael.lawson@newcastle.ac.uk
Dr Maple-Grødem: jodi.maple@uis.no
Prof Tysnes: Ole-Bjorn.Tysnes@uib.no
Dr Williams-Gray: chm27@cam.ac.uk
Dr Yarnall: alison.yarnall@newcastle.ac.uk
Dr Counsell: Carl.Counsell@abdn.ac.uk

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Abstract

Introduction:
Functional dependency in basic activities of daily living (ADLs) is a key outcome in Parkinson’s disease (PD). We aimed to define dependency in PD, using the original and MDS versions of the Unified Parkinson’s Disease Rating Scale (UPDRS).

Methods:
We developed two algorithms to define dependency from items of UPDRS Part 2 and MDS-UPDRS Part 2 relating to basic ADLs (feeding, dressing, hygiene and walking, and getting out of a chair). We validated both algorithms using data from 1110 patients from six community-based PD incidence cohorts, testing concurrent validity, convergent validity, and predictive validity.

Results:
Our optimal algorithm showed high specificity and moderate to high sensitivity versus Schwab & England <80% (specificity 95% [95% confidence interval (CI) 93-97] and sensitivity 65% [95% CI 55-73] at baseline; 88% [95% CI 85-91] and 85% [95% CI 79-97] respectively at five-years follow-up). Convergent validity was demonstrated by strong associations between dependency defined by the algorithm and cognition (MMSE), quality of life (PDQ39), and impairment (UPDRS part 3) (all p <0.001). Algorithm-defined dependency status also predicted mortality: HR for mortality in those dependent vs independent at baseline was 1.6 (95%CI 1.2-2.1) and in those dependent vs independent at five-years’ follow-up was 2.2 (1.6-3.0).

Discussion:
We have demonstrated the concurrent validity, convergent validity, and predictive validity of a UPDRS-/MDS-UPDRS-based algorithm to define functional dependency in PD. This can be used for studying dependency in any study where UPDRS or MDS-UPDRS part 2 data have been collected.
Introduction

Functional dependency is an important patient-orientated outcome in Parkinson’s disease (PD) [1]. Here we focus specifically on dependency in terms of needing help with basic activities of daily living (ADLs), such as washing, dressing, toileting, feeding, or walking. Previous population-based studies have shown that dependency is common, even early in the disease course, [2,3] but there has been little research into the factors that influence this aspect of PD [1].

Existing dependency or activity limitation scales for PD research have drawbacks. The Schwab & England scale has been used most frequently but it is inherently unclear (in particular, the descriptor “chores” is not defined). A Schwab & England score <80% has been used as a definition of dependency (80%=completely independent in most chores; 70%=not completely independent) [3]. The Barthel Index has been used widely in other diseases, but infrequently in PD. Part 2 “Activities of daily living” of the Unified Parkinson’s Disease Rating Scale (UPDRS) and part 2 “Motor Aspects of experiences of daily living” of the Movement Disorders Society (MDS) revision have been used widely, but only a minority of the items in these sections relate to ADLs. If dependency could be identified from the UPDRS part 2, this would allow research into dependency in studies which have collected UPDRS data but not a specific activity limitation/dependency scale. We therefore aimed to develop an algorithm to identify functional dependency using the original UPDRS and MDS-UPDRS Part 2.

Methods

Dataset

We used data from the Parkinson’s Incidence Cohorts Collaboration (PICC), a project to pool data from six PD incidence cohorts in Northern Europe (CamPaIGN4, ICICLE-PD5, NYPUM6, ParkWest7, PICNICS8, and PINE9). These studies each collect demographic, clinical, and genetic data at time of diagnosis and at regular follow-up visits thereafter, with data on 1110 patients at baseline and 714 patients at approximately 5 years. Reasons for missing data at year 5 included (i) death (N=167); (ii) loss to follow-up (N=123); (iii) participants had not reached 5-years’ follow-up or were not seen at about 5 years (N=106). Available data included the original UPDRS (CamPaIGN, NYPUM, ParkWest, PINE), MDS-UPDRS (ICICLE-PD, PICNICS), Schwab & England scale (all studies, except not collected at baseline in PICNICS or ICICLE-PD) and mini-mental state examination (MMSE) (all studies), PDQ-39 (all studies except ParkWest), and Barthel Index (only PINE). Ethical approval for each study was obtained from relevant ethics committees. All participants gave informed consent to participation in the respective studies.

Algorithm development

We developed two algorithms (Table 1) to define dependency using items from UPDRS Part 2 relating to basic ADLs (feeding, dressing, hygiene and walking in both versions and getting out of a chair in MDS-UPDRS). We did not include the turning in bed item from the MDS-UPDRS
because an individual could live independently even if they were unable to turn over in bed. We first assessed what cut-off score for each item would necessarily indicate dependency based on the descriptors, i.e., if the patient had that score or higher they must have been dependent. For instance, for the original UPDRS hygiene item, a score of 3 ("requires assistance for washing, brushing teeth...") was the lowest item which indicated definite evidence of dependency. The cut-offs for each item which indicated patients were clearly dependent are given in table 1. Algorithm 1 was defined as indicating dependency if a patient was clearly dependent in any one of the specified ADL items. Although a simple approach, this algorithm was expected to lead to under-ascertainment of dependency because of the ambiguity in certain descriptors for these items. For example, the original UPDRS hygiene score of 2 ("needs help to shower or bathe; or very slow in hygienic care") could indicate dependency ("needs help to shower or bathe") or independency ("very slow in hygienic care"). Algorithm 2 therefore defined dependency if either (i) any one of these ADL items clearly indicated dependency or (ii) the sum of these ADL items was greater than a certain threshold. To select the threshold we calculated specificity and sensitivity for several thresholds in the PINE study against a “gold standard” of dependency defined by a Barthel Index score of less than 16, excluding the items relating to continence as these are not specific ADLs. We selected the threshold with the maximal accuracy \( \frac{sensitivity + specificity}{2} \). We chose the Barthel Index as a gold standard because a score of less than 16 (after continence items excluded) means the patient needs help with at least one of the following ADLs: mobility, stairs, transfers, dressing, bathing, grooming, feeding, or toilet use. The threshold for the MDS-UPDRS was calculated as the same proportion of maximum score from the specified ADL items.

**Algorithm validation**

For both algorithms, we tested concurrent validity (validating against other instruments that measure the same concept) [10] by assessing specificity and sensitivity of the selected algorithm against dependency defined by Schwab & England scale <80% in each study at baseline (diagnosis) and at five years follow-up (4.5 years in ICICLE-PD). We chose to develop the algorithm 2 cut-off score using the Barthel index in PINE and validate it against the Schwab & England scale in all studies (rather than development using Schwab & England and validation using Barthel in PINE and Schwab & England in the other studies) because i) we wished to use an unambiguous measure of dependency for model development; ii) it allowed all studies to be used in the validation; and iii) it allowed a common and unbiased assessment of heterogeneity across all studies.

We then evaluated convergent validity (evidence of correlation with similar, but not identical, measures) [10] by assessing associations of each algorithm with cognition (MMSE), quality of life (PDQ39), and parkinsonian motor impairment (UPDRS Part 3) at baseline and at 4.5-5 years. For each association, MMSE, PDQ39 or UPDRS part 3 defined the dependent variable
in multivariable linear regression, and the independent variables were dependency as defined by the particular algorithm, age at baseline and sex.

We tested predictive validity by comparing mortality in those defined as dependent and independent by these algorithms in all six studies using Cox regression adjusted for age at baseline and sex. We developed Cox models with survival time measured from (i) baseline and (ii) the 4.5-5 year assessment until death. Patients who had not died were censored at the date last known to be alive.

Statistical analyses were performed using SPSS version Stata version 16 and SPSS version 25.

Results
The PICC cohort included 1110 patients at baseline. Mean age was 69.5 (standard deviation 10.0) and 61% were men. Further characteristics are found in supplementary table 1.

The two algorithms are detailed in table 1. The selected thresholds for algorithm 2 were ≥6 (original UPDRS) and ≥7 (MDS-UPDRS). Concurrent, convergent, and predictive validity are shown in Table 2, Supplementary Tables 2-4, and Supplementary Figures 1-3. The numbers included in each analysis are detailed in the supplementary tables. Algorithm 1 had only low to moderate sensitivities versus Schwab & England (35% [95% CI 26-44] and 58% [95% CI 51-64] at baseline and year 5), but very high specificity (99% [95% CI 98-100] at both time points). Algorithm 2 had higher sensitivities (65% [95% CI 55-73] and 85% [95% CI 79-89] at baseline and year 5), while specificities remained high (95% [95% CI 93-97] and 88% [95% CI 85-91] at baseline and year 5).

There were significant associations between dependency in all the constructs across both algorithms. Using both algorithms, those defined as dependent had significantly higher mortality than those independent at both time points with mortality ratios ranging from 1.6 to 2.6. The results were mostly consistent between studies, but there was some heterogeneity.

Discussion
We have demonstrated the concurrent validity, convergent validity, and predictive validity of two definitions of dependency derived simply from the UPDRS/MDS-UPDRS. Although both algorithms had good face validity, convergent validity and predictive validity, algorithm 2 had better sensitivity while maintaining high specificity. We propose that algorithm 2 is a useful measure of functional dependency in basic ADLs but algorithm 1 may be useful as measure of a more-severe level of dependency. Concurrent validity was better at year 5 than at baseline, probably because baseline dependency relates relatively more to comorbid diseases and later dependency relatively more to PD-related factors; dependency items in UPDRS may be more specific to dependency caused by PD.
This study has several key strengths. The algorithms have been validated in six prospective cohort studies with very low selection bias (each cohort was derived from an incidence study, therefore attempting to include all PD in the population). Therefore, we believe these results are generalizable to Caucasian populations with PD. The data were prospective with a large pooled sample size, and we used multiple methods in our validation.

Nevertheless, this study has some limitations. We lacked a true gold standard for functional dependency. However, we did use validated ADL scales. The Barthel Index has been widely validated in assessing ADLs but has had little validation in PD [11]. Schwab and England has some validation data relating to activity limitation in PD but one study has suggested a score <80% has suboptimal sensitivity as a dichotomous dependency measure [12] despite its face validity. Furthermore, the Barthel data in PINE was self-reported and other scales were reliant on self-report if relatives/carers did not come to study visits so there may be some lack of objectivity. Arguably, the most accurate dependency definition would be to observe patients performing ADLs in their home environment, but this would be resource-intensive and time-consuming and such data were not available. There was heterogeneity in the results between the individual studies included in PICC. There may several reasons for this, including small numbers in individual studies in some of the analyses and variability between studies in how some of the scales were scored. The overwhelming majority of participants were white European so further validation could be done in other ethnic groups.

In conclusion, we have demonstrated the validity of a measure derived from the UPDRS or MDS-UPDRS to simply assess functional dependency in PD. It can therefore be easily applied in most existing clinical studies without additional data collection. Further work is needed to establish which ADL/dependency measure should be the gold standard for use in PD.

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Authors' Roles
Dr Ramsay: 1B, 1C, 2C, 2A, 2B, 3A.
Dr Macleod: 1A, 1B, 1C, 2C, 2A, 2B, 2C, 3B.
Prof Alves: 1A, 1B, 1C, 2C, 3B.
Camacho: 1B, 1C, 2C, 3B.
Prof Forsgren: 1A, 1B, 1C, 2C, 3B.
Dr Lawson: 1A, 1B, 1C, 2C, 3B.
Dr Maple-Grødem: 1B, 1C, 2C, 3B
Prof Tysnes: 1A, 1B, 1C, 2C, 3B.
Dr Williams-Gray: 1A, 1B, 1C, 2C, 3B.
Dr Yarnall: 1A, 1B, 1C, 2C, 3B.
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References


