RESEARCH PAPER

Using ‘dead or dependent’ as an outcome measure in clinical trials in Parkinson’s disease

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ABSTRACT

Background Simple, robust, sensitive and clinically meaningful outcome measures are required for neuroprotective trials in Parkinson’s disease (PD). We explored the feasibility of a composite binary outcome measure, ‘dead or dependent’, in such trials using data from a prospective follow-up study of an incident cohort of PD patients.

Methods Two hundred incident patients had an annual follow-up, including assessment of the Hoehn-Yahr stage (H-Y) and Schwab and England Activities of Daily Living Scale (S&E). Annual scores were converted into binary variables (H-Y <3 vs H-Y ≥3, and S&E ≥80% vs S&E <80%). A new outcome of ‘dead or dependent’ was also created, with dependence in activities of daily living defined as S&E <80%. Using these data, sample sizes were calculated for a hypothetical three-year randomised trial in which the trial outcome was defined by a binary clinical variable, all-cause mortality, or PD-related mortality.

Results At 3 years, 18.0% of patients were dead and 38.4% were dead or dependent. At 80% power, large sample sizes were required if PD-related mortality (n=1938 per study arm) or all-cause mortality (n=734) were used as the outcome, even for large treatment effects (30% reduction in relative risk). The new outcome of ‘death or dependency’ required the smallest sample sizes of all the outcome measures (n=277 for 30% reduction in relative risk, 627 for a 20% reduction).

Conclusions ‘Death or dependency’ is a feasible and potentially useful outcome measure in PD trials of neuroprotective agents, but further work is required to validate its use and define dependency.

INTRODUCTION

Simple, robust and clinically meaningful outcome measures are required for neuroprotective trials in Parkinson’s disease (PD). Clinical outcomes can be classified hierarchically (figure 1) into those which measure impairment (signs of underlying disease, eg, the motor subsection of the old or new Unified Parkinson’s Disease Rating Scale (UPDRS)3), disability (the functional result of impairment, eg, the Schwab and England Activities of Daily Living (ADL) Scale (S&E)3), handicap (the social and societal impact of the disease) and health-related quality of life (eg, the 39-item Parkinson’s disease questionnaire (PDQ-39)). More recently, WHO refined these concepts in terms of impairment in bodily structures/functions, limitations in ADL and restrictions in participation in life situations.4 Measures towards the top of this hierarchy are more likely to be of direct relevance to patients, but are perhaps more susceptible to the confounding effects of factors other than disease progression (eg, mood, finances, comorbidity) than measures towards the bottom of the hierarchy.

In PD neuroprotective clinical trials, disability outcome measures may offer the best compromise between a relevant outcome to patients and as pure as possible a measure of disease progression. Furthermore, disability outcome measures may well be less susceptible to the effects of symptomatic treatment than measures of motor impairment, which are clearly influenced by dopamine replacement therapy.

Loss of independence in either basic ADL (eg, personal hygiene, feeding, ambulation), or instrumental ADL (eg, housework, finances, shopping), are important disability outcomes which should be delayed by neuroprotective treatment. Dependence can be robustly defined, and several valid measures of it exist.3-5 Data relating to dependence also has the advantage of being simple enough to collect via the telephone, helping to minimise missing data in large randomised controlled trials (RCT). However, as with other currently used clinical rating scales, the failure of dependence outcome measures to include ‘death’ as a scale interval necessitates that patients dying during a study must be censored in subsequent analyses, potentially reducing the power of the study to detect a statistically significant improvement in the measure in question.

Censoring in neuroprotective trials would be avoided by using mortality alone as the trial outcome, and this should be reduced/delayed by neuroprotective treatment. However, in diseases such as PD, where patients live for many years after diagnosis, this would require prolonged periods of follow-up and/or very large sample sizes, making such trials largely impractical. A potential solution to these issues may be to use a combined outcome measure which includes mortality and loss of independence in ADL. This approach has already been used in other neurological diseases, such as stroke.6 We, therefore, aimed to explore the feasibility of using a new composite binary outcome measure of ‘dead or dependent’ as opposed to ‘alive and independent’ in PD clinical trials using real-life data from an ongoing prospective follow-up study of a community-based incident cohort of patients with PD.

METHODS

The Parkinsonism Incidence in North-East Scotland (PINE) study is a prospective two-phase incidence
1. Health-related quality of life
2. Handicap / restriction of participation
   • the social impact of disease
3. Disability / limitation in activity
   • the functional result of impairment
4. Impairment of bodily structure / function
   • signs of the underlying disease
5. Pathology
   • Neuronal degeneration / biomarker

**Figure 1** Hierarchy of outcomes measures.

study of parkinsonism in Aberdeen, UK, with preplanned life-
long follow-up of incident patients to establish prognosis. Patients with new-onset possible or probable parkinsonism (defined as two or more of bradykinesia, rest tremor, rigidity or otherwise unexplained postural instability) were identified by a research team from 37 general (primary care) practices (baseline population 315 000) over a four-and-a-half-year period (2002–2004, 2006–2009) using multiple overlapping strategies. Detailed methods and incidence results have been previously published. The study was approved by the Scotland A Multi-centre Research Ethics Committee. All included patients gave their informed consent prior to inclusion in the study.

Ongoing annual follow-up of consenting individuals included assessment of the Hoehn-Yahr (H-Y) stage, and the S&E score. At each follow-up visit, the patient’s clinical diagnosis (PD or an alternative parkinsonian syndrome) was reviewed by a consultant neurologist with an interest in movement disorders, based on factors including the emergence of atypical features (eg, early dementia, falls, marked autonomic features, ataxia, myoclonus, supranuclear gaze palsy), the rate of disease progression, the response to dopamine replacement therapy, the development of motor complications and imaging. The UK PD society brain bank criteria were used to guide the clinical diagnosis of idiopathic PD. During follow-up, patients were invited to consent to postmortem and, if performed, information from these examinations informed the final diagnosis.

Patients could be seen between yearly visits if there was a clinical need, such as to start treatment. Initiation of therapy occurred when a shared decision was made by the patient and doctor. Factors including clinical presentation (tremor less likely to respond than bradykinesia), the impact of symptoms on the patient’s lifestyle and patient preference were taken into consideration, but no specific scores were used to guide initiation of treatment. There was no standard first-line dopaminergic medication; the treatment choice was usually made after discussion of the advantages and disadvantages of each option.

In this report, only consenting patients with a latest or final (if they had died) clinical diagnosis of idiopathic PD were included. Data were extracted from the PINE database on 23 November 2012. For each included patient, data from their baseline appointment, and up to 10 years of annual follow-up data were extracted. No patient had more than 10 years’ follow-up. The date of death for deceased patients was also extracted. Baseline data extracted for each patient included age, gender and the baseline motor UPDRS. For the baseline appointment and each annual follow-up, the H-Y stage and S&E were extracted and converted into binary variables, where people who had died had missing values: H-Y <3 versus H-Y ≥3 (as H-Y ≥3 is the point at which balance disturbance is present); S&E <80% versus S&E ≥80% (as S&E ≤80% is the point at which there is loss of independence in ADL, which in PINE was defined as loss of independence in either washing, dressing, feeding, toileting or mobility). A new binary variable of ‘dead or dependent’ versus ‘alive and independent’ was also computed for each patient at each annual follow-up appointment where, again, dependency was defined by S&E <80%. We also performed a sensitivity analysis where dependency was defined in more severe terms (S&E ≤50% i.e., requiring help with 50% or more of chores).

The PINE study case-notes of included patients were reviewed to determine the type and dose of dopamine replacement therapy taken at each annual follow-up, which were converted into levodopa-equivalent daily doses (LEDD). Study notes, hospital and GP records, and death certificates were reviewed to determine whether those who had died did so as a result of their PD. We considered a ‘PD-related death’ to have occurred when a patient with advanced PD died as the result of a complication (eg, pneumonia, aspiration, infected bed sores, falls, dehydration, or general frailty due to PD). Patients dying suddenly with mild PD, or those with deaths obviously due to another cause (eg, myocardial infarction, stroke, or malignancy), were considered to have had a ‘non-PD related death’.

**Analysis of survival rate**

Using the Kaplan–Meier method, the length of survival from diagnosis (baseline visit) until death was analysed for all included patients. The length of survival in those who had died by the time of data extraction was taken as the difference between the date of death and their baseline (diagnostic) visit, and for those who were still alive, as the difference between the date of data extraction and the baseline visit. A reanalysis of the length of survival, censoring patients with a ‘non-PD related death’ was also undertaken, as was an analysis of time to ‘death or dependency’, whichever occurred first, where dependency was again defined by a S&E <80%.

**Analysis of the change in binary categorical variables over 3 years**

In order to assess whether the binary outcomes changed significantly over time, binary logistic generalised estimating analysis was performed using 3-years of follow-up data from those patients who had completed at least 3 years of follow-up, or would have done so had they not died before the date of data extraction. This time period was chosen, as all except one patient would have completed 3 years’ follow-up when the data were extracted. The change in the binary variables (dead or dependent, S&E <80%, and H-Y ≥3) over 3 years was analysed using data from patients who were free of the respective outcome at baseline. Patients who died were excluded from the H-Y and S&E analyses after the date of death, and the analyses assumed the data were missing completely at random. We, therefore, also did a sensitivity analysis in which those who died were classified as having a poor outcome on the H-Y and S&E. Year of follow-up and gender were included as fixed factors in the model, which was also adjusted for baseline age and motor UPDRS, LEDD at each follow-up and an interaction between LEDD and year of follow-up.

**Sample size calculations for a three-year RCT**

Sample sizes were calculated for a hypothetical three-year parallel-group RCT of a neuroprotective agent in which the trial outcome was either defined by one of the binary variables, all-cause mortality, or PD-related mortality. In these trials, newly diagnosed patients in active or placebo arms would be allowed to commence symptomatic treatment at any time, if required.
However, where one of the binary variables defined the trial outcome, only patients who did not have the respective outcome at baseline would be eligible for inclusion.

Sample sizes were calculated with 80% and 90% power at the 5% level of significance. The proportions of control patients with each outcome were mostly taken from the PINE three-year data to reflect real life. However, a lower level of death or dependency (20%) at three years was also used to reflect what might be seen in milder/younger patients. Although initially the expected treatment effect was taken as a 30% relative risk reduction, this being the level set by the PD-NET collaborative expected treatment effect was taken as a 30% relative risk reduction, this being the level set by the PD-NET collaborative group to detect neuroprotection, a smaller effect (20% relative risk reduction, this being the level set by the PD-NET collaborative group to detect neuroprotection, a smaller effect (20% relative risk reduction) was also assessed to reflect a more modest/realistic treatment effect. Sample sizes (Fisher’s exact method) were calculated using StatsDirect software (V2.7.9). Other statistical analyses were performed using the International Business Machines (IBM) Statistical Package for the Social Sciences (SPSS) Statistics V.20.0.

RESULTS

Of an incident cohort of 377 patients with a baseline diagnosis of possible or probable Parkinsonism, 210 have a latest clinical diagnosis of PD; 10 did not consent to clinical follow-up and were therefore excluded. Of the remaining 200 patients, only one had not completed three years of follow-up by the date of data extraction.

Seventy-one of the 200 included patients died prior to the date of data extraction. Thirty-three died from a PD-related cause (nine from pneumonia; four from dementia; two after a fall; two, who were extremely immobile due to PD, from pulmonary emboli; two with advanced PD from dehydration; two from aspiration; and twelve from general frailty with clearly documented end-stage PD).

Baseline characteristics

The baseline characteristics of the 200 included patients are presented in Table 1. While the majority of patients had mild PD at baseline, around a quarter were already dependent on the help of others for ADL (S&E <80%) at diagnosis, which reflects the elderly nature of the cohort.

### Table 1 Baseline characteristics of the included patients with Parkinson’s disease

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>n</th>
<th>Median value/number with characteristic</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean age in years (SD)</td>
<td>200</td>
<td>72.1 (10.3, range 37–90)</td>
</tr>
<tr>
<td>Male gender (%)</td>
<td>200</td>
<td>118 (59)</td>
</tr>
<tr>
<td>Symptom duration (months)</td>
<td>199</td>
<td>14 (9 to 25, range 2–121)</td>
</tr>
<tr>
<td>UPDRS (I)</td>
<td>166</td>
<td>1 (0 to 2)</td>
</tr>
<tr>
<td>UPDRS (II)</td>
<td>167</td>
<td>10 (6 to 14)</td>
</tr>
<tr>
<td>UPDRS (III)</td>
<td>200</td>
<td>24 (16 to 32)</td>
</tr>
<tr>
<td>Total UPDRS</td>
<td>165</td>
<td>37 (25 to 49)</td>
</tr>
<tr>
<td>H-Y stage ≥3.0 (%)</td>
<td>199</td>
<td>52 (26)</td>
</tr>
<tr>
<td>S&amp;E &lt;80% (%)</td>
<td>200</td>
<td>50 (25)</td>
</tr>
<tr>
<td>GDS-15 ≥5 (%)</td>
<td>156</td>
<td>40 (26)</td>
</tr>
<tr>
<td>MMSE</td>
<td>166</td>
<td>29 (27 to 29)</td>
</tr>
<tr>
<td>Barthel ADL index</td>
<td>158</td>
<td>20 (18 to 20)</td>
</tr>
</tbody>
</table>

*Data are medians with IQRs unless otherwise stated. The ‘n’ column indicates the number of patients who contributed data in each case.*

GDS-15, 15-item Geriatric Depression Scale; H-Y, Hoehn-Yahr stage; MMSE, Mini-mental state examination; S&E, Schwab and England Activities of Daily Living Scale; UPDRS, Unified Parkinson’s Disease Rating Scale.

However, one of the binary variables defined the trial outcome, only patients who did not have the respective outcome at baseline would be eligible for inclusion.

![Figure 2](http://jnnp.bmj.com/)

**Figure 2** Kaplan–Meier curves for different outcomes in PINE study patients with Parkinson’s disease.

**Treatment analysis**

The amount of dopamine replacement therapy taken by patients increased over the 3-year follow-up: median LEDD 0 mg (IQR 0–0) at diagnosis, 240 mg (IQR 0–400) at the first annual follow-up, 300 mg (IQR 100–450) at the second, and 320 mg (IQR 230–500) at the third annual follow-up. Of those who successfully completed the third annual follow up (n=162), 136 (83%) were on dopamine replacement therapy at that time, of whom 24 were on dopamine agonist monotherapy and the remainder levodopa alone or in combination with an agonist or MAO-B inhibitor. Patients not on replacement therapy mostly had milder tremor-dominant disease, and decided against early treatment.

**Survival analysis**

Figure 2 illustrates the Kaplan–Meier survival curves for survival from all-cause mortality, PD-related mortality and time to ‘death or dependence’. The mean survival time from all-cause mortality was shorter (6.7 (95% CI 6.1 to 7.3) years) than that from PD-related mortality (8.1 (95% CI 7.5 to 8.7) years). However, the time until patients reached ‘death or dependence’ was even shorter, with a mean survival time of only 3.6 (95% CI 3.1 to 4.2) years. The stepwise decline seen in the ‘death or dependence’ curve reflects that the S&E, which defined the development of dependency, was mostly collected at each annual follow-up.

**Change in binary categorical variables over 3 years**

Table 2 shows there was a statistically significant increase in the percentage of patients rated negatively in each binary categorical variable over 3 years. The H-Y change became non-significant (p=0.24) in the missing data sensitivity analysis. However, statistically, the most significant change (p=0.027), and the change of greatest magnitude (38.4%), was seen in the composite variable of ‘dead or dependent’ as defined by S&E <80%.

**Sample size calculations**

Table 3 shows that using either all-cause mortality or PD-related mortality as the trial outcome for the three-year RCT would
DISCUSSION

Our data show that a composite binary measure of ‘death or dependency’ as opposed to ‘alive and independent’ is potentially a simple and feasible new outcome measure for clinical trials in PD.

Neuroprotective trials in PD have struggled to separate out symptomatic effects of potential therapeutic agents from true disease-modifying effects. Limitations in trial designs aimed at detecting a neuroprotective effect (eg, the measurement of clinical outcomes following a wash-out period) and lower rates of outcomes events (20%) were expected (table 3). Given that it might be argued that combining death and mild degrees of dependency is inappropriate, the sample size for detecting death or more major dependency (S&E ≤ 50%) was also calculated using the number who were dead or had S&E ≤ 50% at 3 years. Significantly higher sample sizes were again required than when the higher cut-off value (S&E < 80%) was used (table 3).

However, long-term follow-up studies have several disadvantages. They are expensive and may take many years until results are available. Furthermore, high drop-out rates may be encountered, particularly if the coadministration of symptomatic treatment is prohibited, which may make interpretation of results difficult or impossible. For example, after 7 years of follow-up in a study comparing clinical outcomes in patients treated with selegiline versus those treated with placebo, only 35% of participants initially randomised to treatment were still in the study.16 Several of these problems would be overcome in a three-year trial that used ‘death or dependency’ as the primary outcome. First, symptomatic treatment would be permitted in addition to the study drug as, even on such treatment, we have shown significant numbers of patients develop the outcome of interest. This will also limit drop-outs. Secondly, by limiting follow-up to three years, we have hopefully struck a balance between a significantly long enough period of follow-up to detect a neuroprotective effect, and a realistic period of follow-up, financially, and in terms of patient compliance.

Our new composite measure of ‘death or dependency’ using a S&E cut-off of 80% has the advantage of being more clinically relevant to patients than a change in impairment measured by the motor UPDRS, and would be expected to change in response to an effective neuroprotective drug. It also avoids the need to censor data from patients who die during a study, thereby maximising the power of that study. As a clinical outcome measure, we have demonstrated that it is sensitive to disease progression even with the coadministration of symptomatic dopamine replacement therapy. Our sample size calculations demonstrated that a three-year RCT using this composite measure would be feasible, requiring about 600–800 patients per arm, depending on power, assuming a more realistic but still clinically valuable 20% relative risk reduction, and the same rate of death or dependency (38%) as we found at 3 years. Even if

Table 2 Percentage of patients with each binary outcome at each annual follow-up over three years

<table>
<thead>
<tr>
<th>Binary outcome category</th>
<th>Baseline n (%)</th>
<th>Year 1 n (%)</th>
<th>Year 2 n (%)</th>
<th>Year 3 n (%)</th>
<th>Overall p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>H-Y stage ≥ 3</td>
<td>0/147 (0)</td>
<td>16/132 (12.1)</td>
<td>30/134 (22.4)</td>
<td>28/127 (22.0)</td>
<td>0.047</td>
</tr>
<tr>
<td>S&amp;E &lt;80%</td>
<td>0/151 (0)</td>
<td>21/144 (14.6)</td>
<td>34/136 (25.0)</td>
<td>37/130 (28.5)</td>
<td>0.045</td>
</tr>
<tr>
<td>‘Dead or dependent’</td>
<td>0/151 (0)</td>
<td>28/151 (18.5)</td>
<td>48/150 (32.0)</td>
<td>58/151 (38.4)</td>
<td>0.027</td>
</tr>
</tbody>
</table>

The ‘n’ column indicates the number of patients in the stated binary category compared with the total number of patients who contributed data relevant to that binary variable at each point in time. The % column indicates the actual (not modelled) percentage of patients in the stated binary category at each annual follow-up. The overall P values, derived from generalised estimating equations, relate to the significance of the change in each variable over three years.

H-Y, Hoehn-Yahr scale; S&E, Schwab and England Activities of Daily Living Scale.

Table 3 Sample sizes required to appropriately power RCTs of three years duration in patients with idiopathic PD

<table>
<thead>
<tr>
<th>Primary outcome measure</th>
<th>Event rate in control arm (%)</th>
<th>Relative risk reduction (%)</th>
<th>Absolute risk reduction (%)</th>
<th>Sample size in each study arm (80% power)</th>
<th>Sample size in each study arm (90% power)</th>
</tr>
</thead>
<tbody>
<tr>
<td>PD-related mortality</td>
<td>7.5</td>
<td>30</td>
<td>2.25</td>
<td>1938</td>
<td>2564</td>
</tr>
<tr>
<td>All-cause mortality</td>
<td>18.0</td>
<td>30</td>
<td>5.40</td>
<td>734</td>
<td>969</td>
</tr>
<tr>
<td>H-Y stage ≥ 3</td>
<td>22.0</td>
<td>30</td>
<td>6.60</td>
<td>577</td>
<td>762</td>
</tr>
<tr>
<td>S&amp;E &lt;80%</td>
<td>28.5</td>
<td>30</td>
<td>8.55</td>
<td>417</td>
<td>549</td>
</tr>
<tr>
<td>Dead or dependent (S&amp;E &lt;80%)</td>
<td>38.4</td>
<td>30</td>
<td>11.52</td>
<td>277</td>
<td>364</td>
</tr>
<tr>
<td>Dead or dependent (S&amp;E &lt;80%)</td>
<td>38.4</td>
<td>20</td>
<td>7.68</td>
<td>627</td>
<td>830</td>
</tr>
<tr>
<td>Dead or dependent (S&amp;E ≤50%)</td>
<td>20.0</td>
<td>20</td>
<td>4.00</td>
<td>1497</td>
<td>1987</td>
</tr>
<tr>
<td>Dead or dependent (S&amp;E ≤50%)</td>
<td>22.6</td>
<td>30</td>
<td>6.78</td>
<td>559</td>
<td>737</td>
</tr>
</tbody>
</table>

Sample sizes were calculated with a 5% level of significance.

H-Y, Hoehn-Yahr scale; S&E, Schwab and England Activities of Daily Living Scale.
very large trials were required (eg, 1500–2000 per arm to
detect, to a reduction in poor outcome from 20% to 16%), the
simplicity and ease of collection of the measure would also
allow it to be used in simple pragmatic trials, perhaps using
large prospective patient registries.19

There are a number of potential limitations in our trial
design. This was a small study and needs validation in larger
cohorts. Patients also needed to be independent in ADL at base-
line, which meant 25% were excluded (this proportion would
be lower in younger cohorts). However, this is already the case
in neuroprotection trials where patients usually have early stage
disease, implying a reasonable proportion of intact neurons
remain which may be salvageable. However, if required, an
adaptation in the outcome would also allow dependent patients
to be included, that is, including deterioration to a higher grade
of dependency as an outcome in those already dependent. This
has been used in stroke trials.

Our data come from a community-based incident cohort of
PD. While this makes it more generalisable to the totality of PD
patients in the community, it will be less applicable to hospital-
based cohorts of people with early PD, who are the types of
patient recruited into RCTs. In particular, our patients were
older than most patients recruited into RCTs.15 The proportion
of patients dead or dependent at 3 years will vary by age of
onset, and so much larger (or longer) trials would be required if
younger patients were recruited. For example, we showed that
reducing the outcome event rate to 20% at 3 years would
require very large trials to detect 20% relative risk reductions.
However, recruitment of older people into PD trials should be
encouraged to improve generalisability and clinical relevance,
given that PD is predominantly a disease of the elderly.

Some of the outcomes (deaths and loss of independence in
ADL) were the result of comorbid disease, not PD, and so
would not be expected to be prevented by neuroprotective treat-
ments. Restricting outcomes to those thought to be due solely
to PD might seem sensible, but would add a degree of subjectiv-
ity (particularly when deciding about loss of independence) and
would also dramatically increase sample sizes as shown by the
data for mortality alone. Moreover, trials using impairment
scores, such as the motor UPDRS, are also faced with a similar
problem as it remains unclear how to score these in the presence
of significant comorbidity.19

Finally, it might be argued that combining death and any
dependency in ADL is inappropriate because the two are quite
different. However, this outcome is already widely used in major
stroke trials8 and meta-analyses,20 and other fields of medicine
frequently use endpoints composed of fatal and non-fatal out-
comes, for example, cardiovascular trials often combine cardio-
vascular death along with non-fatal myocardial infarction and
stroke. Any effective neuroprotective agent would be expected to
reduce PD mortality and delay progression to dependency in sur-
vivors, so clinically it makes sense to combine the two and, as
highlighted already, it allows all patients randomised to be
included in the analyses at every time point. Using more severe
dependency (eg, S&E \( \leq 50 \)) in combination with death can be
(done and again is used in stroke trials to measure ‘devastating
outcome’),21 but we showed it does increase the sample size sig-
nificantly, even for large treatment effects (table 3).

Although ‘death or dependency’ looks promising as an
outcome, further work is required before it can be widely
adopted. In particular, although we used the S&E scale and
clearly defined what we regarded as becoming dependent
(\(<80\%\)) in the PINE study, the overall validity and reproducibil-
ity of the S&E scale has been poorly studied. There are many
other disability scales that could be used, which may be better.
However, we would caution against the use of the HY scale
because it mixes impairment and disability, and there is no clear-
cut point where someone becomes dependent. It would also be
crucial to define dependency more clearly. For example, depend-
ence in instrumental ADL is different to dependency in basic
ADL, and may be more sensitive to early change in PD.

In conclusion, we believe that ‘death or dependency’ is a
potentially useful new simple outcome measure for PD trials,
particularly neuroprotective trials, but more work needs to be
done to confirm this.

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Data sharing statement The PINE data are available for sharing by approaching
Dr Counsell.

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