Performance on the dementia rating scale in Parkinson’s disease with dementia and dementia with Lewy bodies: comparison with progressive supranuclear palsy and Alzheimer’s disease

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Background: The relation between dementia with Lewy bodies (DLB) and Parkinson’s disease with dementia (PDD) is unknown.

Objectives: To compare the cognitive profiles of patients with DLB and PDD, and compare those with the performance of patients with a subcortical dementia (progressive supranuclear palsy) and a cortical dementia (Alzheimer’s disease).

Design: Survey of cognitive features.

Setting: General community in Rogaland county, Norway, and a university dementia and movement disorder research centre in the USA.

Patients: 60 patients with DLB, 35 with PDD, 49 with progressive supranuclear palsy, and 29 with Alzheimer’s disease, diagnosed by either standardised clinical procedures and criteria (all PDD and Alzheimer cases and 76% of cases of progressive supranuclear palsy), or necropsy (all DLB cases and 24% of cases of progressive supranuclear palsy). Level of dementia severity was matched using the total score on the dementia rating scale adjusted for age and education.

Main outcome measures: Dementia rating scale subscores corrected for age.

Results: No significant differences between the dementia rating scale subscores in the PDD and DLB groups were found in the severely demented patients; in patients with mild to moderate dementia the conceptualisation subscore was higher in PDD than in DLB (p = 0.03). Compared with Alzheimer’s disease, PDD and DLB had higher memory subscores (p < 0.001) but lower initiation and perseveration (p = 0.008 and p = 0.021) and construction subscores (p = 0.009 and p = 0.001). DLB patients had a lower conceptualisation subscore (p = 0.004). Compared with progressive supranuclear palsy, PDD and DLB patients had lower memory subscores (p < 0.001).

Conclusions: The cognitive profiles of patients with DLB and PDD were similar, but they differed from those of patients with Alzheimer’s disease and progressive supranuclear palsy. The cognitive pattern in DLB and PDD probably reflects the superimposition of subcortical deficits upon deficits typically associated with Alzheimer’s disease.

Dementia with Lewy bodies (DLB) and Parkinson’s disease with dementia (PDD) are common syndromes characterised by parkinsonism and dementia. There are few comparisons of the clinical presentations of these syndromes, and differentiation between the disorders may occasionally be difficult in clinical practice. Moreover, although there is some evidence suggesting pathological, neurochemical, and clinical similarities, the nosological relation between the two syndromes has yet to be resolved.

The cognitive impairment in DLB is characterised by marked deficits in attention and in executive, visuospatial, and constructional abilities. Memory is also impaired, although recall is relatively spared when compared with Alzheimer’s disease. A similar pattern of cognitive impairment is found in Parkinson’s disease. Nevertheless, two studies that directly compared cognition in PDD and DLB reported more executive dysfunction in the latter. However, the mean mini-mental state examination (MMSE) score was only 12.5 in the DLB group compared with a nearly normal score of 24.1 in the Parkinson’s disease group.

To ensure that any between-group differences in subtest profiles are attributable to differences in the underlying nature of the two diseases rather than to differences in the global level of dementia, the two groups should be matched for overall severity of dementia. In the study by Downes et al., clinically diagnosed PDD and DLB cases were carefully matched for dementia severity, but the study sample was small (only 10 patients in each group), the subjects were rather young (mean ages 64.9 and 66.4 years, respectively), and the level of cognitive impairment was mild (mean verbal IQ 97.2 and 94.1, respectively). Accordingly, the representativeness of these samples can be questioned.

Cortical and subcortical pathologies occur in both DLB and PDD, with potential implications for cognitive function, but the brain structures contributing to the cognitive impairment are still not clearly identified. Furthermore, differentiation of patients with DLB or PDD from those with dementia syndromes characterised by symptoms of mainly cortical pathology, such as Alzheimer’s disease, or subcortical pathology, such as progressive supranuclear palsy, may be difficult. Knowledge of the cognitive patterns in DLB and PDD in comparison with Alzheimer’s disease and progressive supranuclear palsy could provide information about factors contributing to the cognitive

Abbreviations: DLB, dementia with Lewy bodies; MMSE, mini-mental state examination; NINDS, National Institutes of Neurological Disorders and Stroke; PDD, Parkinson’s disease with dementia; SPSP, Society for Progressive Supranuclear Palsy

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dysfunction in these disorders, and could aid in the clinical diagnosis of dementia with parkinsonian syndromes and provide information on the nosological status of PDD and DLB.

With these aspects in mind, we used the dementia rating scale—a brief measure of general cognitive status commonly used in both clinical practice and research—to study the performance of relatively large samples of patients with PDD, DLB, progressive supranuclear palsy, and Alzheimer’s disease with similar levels of overall dementia. We hypothesised that the cognitive profiles of PDD and DLB cases are rather similar and are consistent with the presence of both cortical and subcortical dysfunction, in contrast with syndromes of cortical dementia (Alzheimer’s disease) and subcortical dementia (progressive supranuclear palsy).

**METHODS**

**Subjects**

**Parkinson’s disease**

PDD patients were recruited from a community based cohort of subjects with Parkinson’s disease in the county of Rogaland, Norway. They were evaluated using a comprehensive battery of neurological, psychiatric, and neuropsychological tests at four year intervals for eight years. The cohort was acquired after a detailed search, and probably represents the entire Parkinson’s disease population in the region (see Tandberg et al for details). Patients who developed dementia more than one year after the onset of parkinsonism were included. In order to exclude DLB from this group, patients with dementia, repeated falls, or hallucinations at disease onset, or dementia at the primary evaluation, were not included.

A diagnosis of clinical definite, probable, or possible Parkinson’s disease was made according to previously published criteria. In order to obtain a homogeneous population with a high diagnostic specificity for Parkinson’s disease, only subjects with definite Parkinson’s disease were included in the study. This required that a patient had resting tremor and at least two of the three other cardinal signs of akinesia, rigidity, and postural abnormalities. Unilateral onset and development of parkinsonism as well as a good to excellent response to a dopaminergic agent were also required.

Atypical features such as early falls, autonomic failure, or dementia, and pyramidal signs, cerebellar signs, or supranuclear gaze palsy on examination excluded the diagnosis of clinically definite Parkinson’s disease. Additional exclusion criteria were other diseases that could explain the symptoms. Radiological structural brain abnormalities not compatible with a diagnosis of Parkinson’s disease, a history of alcohol or substance abuse in the past year, head trauma with loss of consciousness, and psychiatric disorders other than depression requiring treatment preceding the onset of the current disease.

**Dementia with Lewy bodies**

We included all 62 patients with necropsy confirmed DLB from the Alzheimer Research Center in San Diego (University of California San Diego) in whom the dementia rating scale had been administered during life. All these were diagnosed as having dementia at the time of their initial evaluation. They had received yearly physical, neurological, and neuropsychological assessments. Informed consent for necropsy was obtained from the next of kin immediately after the patient’s death. Twenty three of the patients have been reported previously. The procedures that were used to prepare and study the brains of the DLB patients have been described in detail before. The neuropathological diagnosis of DLB required the presence of subcortical and cortical Lewy bodies. The majority of the DLB subjects also had some degree of neocortical senile plaque formation.

**Progressive supranuclear palsy**

The patients with progressive supranuclear palsy were 55 consecutive outpatients with this condition presenting to the National Institutes of Neurological Disorders and Stroke (NINDS), Bethesda, Maryland, USA, for evaluation and participation in research studies, and who fulfilled the research criteria of the NINDS-SPSP (Society for Progressive Supranuclear Palsy) for the diagnosis of definite (n = 13), probable (n = 34), or possible (n = 8) progressive supranuclear palsy. Definite progressive supranuclear palsy required eventual neuropathological confirmation. The reliability and validity of the “possible” and “probable” NINDS-SPSP criteria for the diagnosis of this condition are high (as confirmed by pathology). This is particularly the case for “probable” diagnosis (positive predictive value and specificity, 100%).

**Alzheimer’s disease**

We analysed the dementia rating scale scores from 39 consecutive patients with Alzheimer’s disease attending the section of geriatric psychiatry, Rogaland Psychiatric Hospital, Stavanger, Norway. They had no clinically significant psychiatric symptoms. They were diagnosed according to ICD-10 criteria for Alzheimer’s disease. The diagnosis was based on interviews with patients and caregivers, cognitive testing, physical examination, routine blood tests, and cranial computed tomography. None of the patients had parkinsonism.

**Procedures**

All patients, and when appropriate their carers, gave informed consent for participation in this research study. The dementia rating scale and the MMSE were administered according to standardised instructions. The dementia rating scale is divided into five subtests, measuring attention, initiation and perseveration, construction, conceptualisation, and memory. For individuals with Parkinson’s disease, the scale has been shown to be a valid and screening test of cognitive functioning, and the subtests show strong convergent and discriminant validity. To obtain the maximum number of patients with dementia, we used the most recent evaluation of the patients with Parkinson’s disease. Those with motor fluctuations were assessed in the “ON” phase. The stage of parkinsonism in the patients with Parkinson’s disease and progressive supranuclear palsy was rated by a neurologist using the Hoehn and Yahr scale. The presence of depression was determined by a separate score of 3 or more on the neuropsychiatric inventory depression item in Parkinson’s disease, progressive supranuclear palsy, and Alzheimer’s disease, and by the diagnostic interview schedule in the DLB group. Although the neuropsychiatric inventory is caregiver based, it significantly correlates with standard patient based rating scales of depression.

**Statistical analysis**

Performance on the dementia rating scale is sensitive to age and education, but not sex. To adjust for differences in age and education in the groups, the total score on the dementia rating scale was converted to an age and education adjusted dementia rating scale subscores—were analysed to adjust for differences in age and education in the groups, the total score on the dementia rating scale was converted to an age and education adjusted dementia rating scale subscores—were analysed on August 5, 2023 by guest. Protected by copyright.
RESULTS

Thirty five cases of PDD, 60 of DLB, 49 of progressive supranuclear palsy, and 29 of Alzheimer’s disease had mild to moderate or severe dementia; thus the study sample consisted of 173 subjects. The mean (SD) duration of parkinsonism preceding the onset of dementia in the PDD group was 12.3 (4.8) years. The mean Hoehn and Yahr stage was 3.5 (0.9) in the patients with Parkinson’s disease and 3.8 (0.7) in the patients with progressive supranuclear palsy (NS). Other clinical and demographic characteristics are shown in table 1. The groups differed with regard to sex, age, years of education, MMSE, and total score on the raw dementia rating scale. There were mostly small, but statistically significant, differences in the age and education corrected total scores on the dementia rating scale in the groups with mild to moderate dementia (F = 14.6; df = 3,53; p < 0.001) and severe dementia (F = 13.3; df = 3,112; p < 0.001).

The pairwise comparisons showed that in the mild to moderately demented cohort, the age and education corrected total score on the dementia rating scale in the Alzheimer group was significantly higher than in the PDD and DLB groups, while the other groups did not differ substantially; in the severely demented cohort, the score in the PDD group was significantly higher than in the DLB and progressive supranuclear palsy groups, and higher in the Alzheimer group than in the DLB group.

Depression was diagnosed in five patients with PDD (18%), in nine with DLB (15%), in nine with progressive supranuclear palsy (18%), but in none of the Alzheimer cases (p = 0.115). All patients with PDD, 10 patients with progressive supranuclear palsy (20%), and two patients with DLB (3%) were taking antiparkinsonian agents. Psychotropic drugs with potentially adverse effects on cognition (tricyclic antidepressants, benzodiazepines, traditional antipsychotics) were being used by seven patients with PDD (20%), six with DLB (10%), and six with progressive supranuclear palsy (12%), but by none of the patients with Alzheimer’s disease (p = 0.085). There were no significant associations between age and education adjusted total scores on the dementia rating scale and the presence or severity of depression or the use of psychotropic agents in any of the patient groups.

In the mild to moderate dementia cohort, the one way ANOVA showed highly significant between-group differences for the age adjusted dementia rating scale subscores “initiation and perseveration” (F = 6.7; p = 0.001), “construction” (F = 5.5; p = 0.002), “conceptualisation” (F = 6.9; p = 0.001), and “memory” (F = 16.3; df = 2,43; p < 0.001), but not for “attention” (F = 0.7, p = 0.57) (df = 3,53 for all analyses except “memory”). In the cohort with severe dementia, the groups differed significantly on the subscores for “construction” (F = 8.9; p < 0.001) and “memory” (F = 32.3; df = 2,43; p < 0.001). As seen in figs 1 and 2 it was mainly the progressive supranuclear palsy group that differed from the others, with higher “memory” scores than the PDD and DLB groups (p < 0.001). The “memory” scores of the PDD and DLB groups were not significantly different either for the mild to moderate or the severe dementia range. The “memory” scores were lower in the Alzheimer group than in the PDD and DLB groups with either mild to moderate dementia (p = 0.002) or severe dementia (p = 0.018). The proportion of patients with the lowest possible score (2) was significantly higher in the Alzheimer group than in the PDD group (p = 0.0007 for mild to moderate dementia and p = 0.018 for severe dementia).

### Table 1 Characteristics of the patients

<table>
<thead>
<tr>
<th></th>
<th>Mild to moderate dementia</th>
<th>Severe dementia</th>
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<tbody>
<tr>
<td></td>
<td>PDD (n=17)</td>
<td>DLB (n=9)</td>
</tr>
<tr>
<td>Age at testing (years)</td>
<td>77.2 (5.6)</td>
<td>73.2 (7.2)</td>
</tr>
<tr>
<td>Female (%)</td>
<td>24</td>
<td>56</td>
</tr>
<tr>
<td>Duration of disease (years)</td>
<td>15.4 (5.0)</td>
<td>1.9 (2.1)**</td>
</tr>
<tr>
<td>Education (years)</td>
<td>9.5 (4.1)</td>
<td>12.0 (4.6)</td>
</tr>
<tr>
<td>MMSE score</td>
<td>23.6 (2.6)</td>
<td>23.9 (2.8)</td>
</tr>
<tr>
<td>DRS score, uncorrected</td>
<td>120.6 (7.0)</td>
<td>118 (10.8)</td>
</tr>
<tr>
<td>DRS scores, corrected for age and education</td>
<td>48.1 (1.4)</td>
<td>3.9 (1.2)</td>
</tr>
</tbody>
</table>

Values are mean (SD).

*The diagnostic groups were compared separately in the mild and moderate-severe dementia groups using one way analysis of variance. Scheffe’s post hoc test was used for pairwise comparisons. Sex distribution was compared using χ² tests.

Different from PDD: *p < 0.05, **p < 0.001; different from DLB: †p < 0.05, ††p < 0.001.

AD, Alzheimer’s disease; DRS, dementia rating scale; DLB, dementia with Lewy bodies; MMSE, mini-mental state examination; PDD, Parkinson’s disease with dementia; PSP, progressive supranuclear palsy.
The proportion of patients with the lowest possible score was higher in the Alzheimer group than in the group with progressive supranuclear palsy ($p = 0.0001$ in both severity groups).

DISCUSSION

Our main finding was a similar pattern of cognitive impairment in patients with PDD and DLB who had severe dementia, and on four of the five subscores in those with mild to moderate dementia. This finding extends recent comparative studies of motor, psychiatric, and attentional disturbances in PDD and DLB, highlighting the similarities between these syndromes. Given the neurochemical (for example, the cholinergic deficits) and pathological (cortical Lewy bodies) similarities in PDD and DLB, the clinical similarities are not surprising. The current distinction between these two conditions is based upon the timing of symptom onset. Patients are classified as having DLB when cognitive impairment or hallucinations present before or within one year of the onset of dementia.
year of the onset of parkinsonism; they are classified as having PDD when the parkinsonism precedes the dementia by more than one year. On the basis of recent findings, PDD and DLB might better be viewed as parts of a spectrum of Lewy body disorders.

On the other hand, as both our study and a previous one show that executive function in patients with mild dementia and DLB is more impaired than in patients with mild PDD, it is hypothesised that the involvement of the frontal cortex occurs later in PDD. The dementia rating scale does not permit further evaluation of what leads to the memory disturbance in these two disorders, as we cannot determine whether patients fail to encode or to retrieve information. This would be necessary to establish whether the deficit results from lesions in the hippocampus or in the frontal cortex. Alternatively, it is possible that the between-group minor differences at early disease stages may reflect the heterogeneity of the underlying lesions in PDD and DLB (Lewy bodies, amyloid plaques, parkinsonism associated with other disorders, vascular disease).

We found that the cognitive pattern of patients with DLB and PDD differed from that of patients with cortical dementia (Alzheimer's disease) and subcortical dementia (progressive supranuclear palsy). On the memory subscale of the dementia rating scale—which is related to hippocampal atrophy—the DLB and PDD subjects were more impaired than the patients with progressive supranuclear palsy, and less impaired than the patients with Alzheimer's disease. These findings are consistent with previous reports. 15 19 20 Hippocampal atrophy is found in Parkinson's disease and DLB, and is more pronounced in DLB than in progressive supranuclear palsy. 29 The “initiation and perseveration” subscore, related to frontal atrophy, and the “construction” score, related to bilateral temporal atrophy, were less impaired in Alzheimer's disease than in PDD and DLB, consistent with previous studies. 29 20 The cognitive pattern in patients with DLB and PDD may be characterised as reflecting the superimposition of fronto-subcortical cognitive deficits upon the cognitive deficits typically associated with Alzheimer's disease. A confounding factor in this study was that diagnostic group was segregated by research centre. Thus the findings may have been influenced by different approaches to diagnosis or evaluation. However, the clinical diagnoses were based on standardised procedures and criteria, and the diagnosis of all DLB patients and of at least 85% of patients with progressive supranuclear palsy was either confirmed by necropsy using standardised criteria (defined as 100% accuracy) or met the diagnostic criteria. The PDD patients were followed longitudinally by the same neurologists for at least four years, and all DLB patients and of at least 85% of patients with progressive supranuclear palsy were followed by the same neurologists for at least four years. Thus, the dementia rating scale— which is related to hippocampal atrophy—is the minimally affected in the four groups. How- ever, these differences may have influenced the pattern of cognition differently in the four groups. 33

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Furthermore, our groups differed with regard to sample selection. The PDD patients were drawn from an epidemiological population of cases of Parkinson's disease and therefore represent the full spectrum of disease severity and presentation. The DLB and progressive supranuclear palsy groups, on the other hand, were referrals to university dementia or movement disorder research centres, introducing a potential bias towards the inclusion of more severely affected or complicated cases and the exclusion of mild cases. The groups differed with regard to age, sex, and years of education, and their performance on the dementia rating scale is influenced by age and the level of education, but not by sex. 24 27 To control for potential bias based on demographic differences, the overall level of dementia was based on age and education corrected norms, and the dementia rating scale subscores were converted to age corrected scores. An education based differential influence on the dementia rating scale performance cannot be entirely excluded, however. The strengths of our study include the relatively large samples, the prospective design that aided in the diagnosis of PDD, and the necropsy confirmation in DLB—the most difficult of the syndromes to diagnose clinically—and in 24% of the patients with progressive supranuclear palsy. The DLB and progressive supranuclear palsy groups are among the largest cohorts yet reported, and the PDD group was community based. Thus the patient groups may be representative of patients with these brain diseases. The overall severity of dementia was comparable in the four diagnostic groups. However, there were small between-group differences, particularly in patients with severe dementia. It is possible that these differences may have influenced the pattern of cognition differentially in the four groups.

Conclusions

A similar pattern of cognitive impairment was found in patients with PDD and DLB with severe dementia, and was different from that in patients with cortical and subcortical dementia. Our findings suggest but do not prove that the underlying mechanisms of cognitive decline in Parkinson's disease are similar to those in DLB, but different from those in Alzheimer's disease and progressive supranuclear palsy.
REFERENCES


