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EXPERIENCE

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ABC Corporation - JUNE 2013 - OCTOBER 2013

- Philippines Emphasized critical thinking, basic grammar, and composition skills for effective writing.
- Administered and graded tests and assignments to evaluate student progress.
- Assigned lessons and corrected homework.
- Improved instruction methods by using a variety of assessment tools and strategies.
- Worked with an average of 45 students per class.
- Advised students in the thesis writing process.
- Served as Thesis panel member and Chair for student defenses.

University Professor

Delta Corporation - 2008 - 2013

- Business, marketing, and tourism programs.
- Content development concerning transportation systems and material storage, aimed at reducing costs in the class of Material Management (7th Semester).
- Created and designed clothes and accessories Designed environments, costumes, and lighting for theater productions, film, opera, ballet, and others.
- Planned and developed displays in-store.
- Employed by both ground and online universities.
- Taught multiple courses in the catalog both under the psychology core courses as well as the substance.
- Personality, Personality theories, Motivation, Cognitive, Child Development, Intro, and Advanced Counseling, Substance Abuse.

EDUCATION

- Master Of Arts

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Translational
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RESEARCH

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Caregiver burden is increased in Parkinson's disease with mild cognitive impairment (PD-MCI)

Ann J. Jones^{1,2*}, Roeline G. Kuijer¹, Leslie Livingston^{1,3}, Daniel Myall^{1,3}, Kyla Horne^{1,2,3}, Michael MacAskill^{1,3}, Toni Pitcher^{1,3}, Paul T. Barrett^{1,4}, Tim J. Anderson^{1,3,5,6} and John C. Dalrymple-Alford^{1,2,3,6}

Abstract

Background: There is limited evidence on caregiver outcomes associated with mild cognitive impairment in patients with Parkinson's disease (PD-MCI) and the coping strategies used by these caregivers.

Methods: To investigate this relationship, we examined levels of burden, depression, anxiety, coping strategies and positive aspects of caregiving in the informal caregivers of 96 PD patients. The PD patients were classified using MDS-Task Force Level II criteria as showing either normal cognition (PD-N; n = 51), PD-MCI (n = 30) or dementia (PDD; n = 15).

Results: Mean Zani Burden Interview (ZBI) score increased significantly between carers of PD-N ($M = 13.39$, $SD = 12.22$) compared to those of PD-MCI patients ($M = 22.00$, $SD = 10.8$), and between carers of PD-MCI and PDD patients ($M = 29.33$, $SD = 9.59$). Moreover, the proportion of carers showing clinically significant levels of burden (ZBI score ≥ 21) also increased as the patients' cognitive status declined (18% for PD-N; 60% for PD-MCI and 80% for PDD) and was mirrored by an increasing amount of time spent providing care by the caregivers. Caregiver ZBI score was independent of patient neuropsychiatric symptoms, motor function, disease duration and time that caregivers spent caregiving. Caregiver use of different coping strategies increased with worsening cognition. However, we found only equivocal evidence that the use of problem-focused, emotion-focused and dysfunctional coping mediated the association between patient cognitive status and caregiver burden, because the inverse models that used caregiver burden as the mediator were also significant.

Conclusions: The study highlights the impact of Parkinson's disease on those providing care when the patient's cognition is poor, including those with MCI. Caregiver well-being has important implications for caregiver support, nursing home placement and disease course.

Keywords: Parkinson's disease, Mild cognitive impairment, Zani caregiver burden interview, Coping, Depression, Anxiety, Positive aspects of caregiving

Background

Cognitive problems are an integral part of Parkinson's disease because dementia (PDD) eventuates in 75%–90% of patients and is associated with substantial health and economic burden [1]. Patients who present with mild cognitive impairment (PD-MCI) are at high risk of

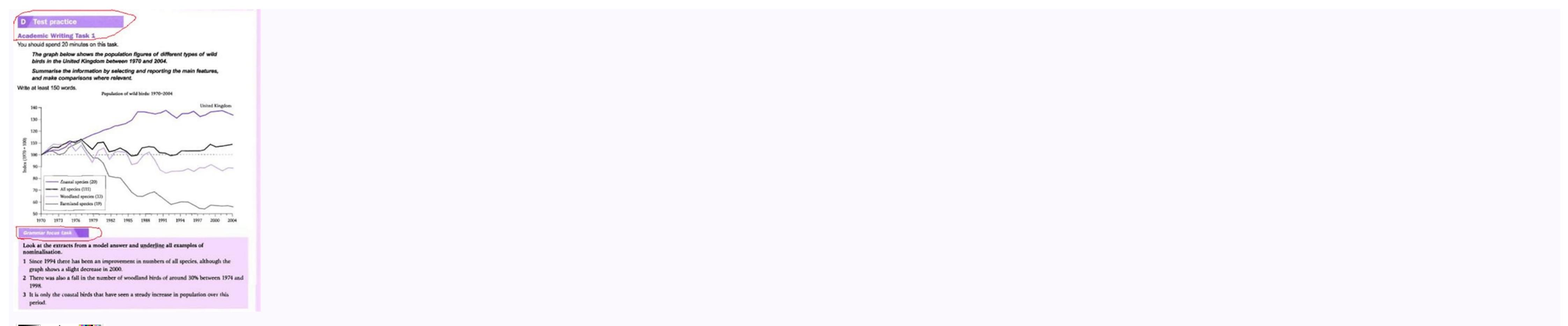
progression to PDD and are a growing research focus to assess the impact of increasing cognitive impairment and to facilitate early intervention [2–5]. The impact of cognitive status on the informal caregiver is a relatively neglected dimension of PD-MCI and there is limited research on the specific effects of cognitive decline on the well-being of caregivers of these PD patients. Understanding these relationships is important because improving caregiver support and well-being may have a bearing on disease management and delay formal care and nursing home placement.

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