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Predictors of agreement between general practitioner detection of dementia and the revised Cambridge Cognitive Assessment (CAMCOG-R)

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ABSTRACT

Background: Dementia is a complex and variable condition which makes recognition of it particularly difficult in a low prevalence primary care setting. This study examined the factors associated with agreement between an objective measure of cognitive function (CAMCOG-R) and general practitioner (GP) clinical judgment of dementia.

Methods: This was a cross-sectional study involving 165 GPs and 2,024 community-dwelling patients aged 75 years or over. GPs provided their clinical judgment in relation to each of their patient's dementia status. Each patient's cognitive function and depression status was measured by a research nurse using the CAMCOG-R and the 15-item Geriatric Depression Scale (GDS) respectively.

Results: GPs correctly identified 44.5% of patients with CAMCOG-R dementia and 90% of patients without CAMCOG-R dementia. In those patients with CAMCOG-R dementia, two patient dependent factors were most important for predicting agreement between the CAMCOG-R and GP judgment: the CAMCOG-R score ($p=0.006$) and patient's mention of subjective memory complaints to the GP ($p=0.040$). A higher CAMCOG-R ($p<0.001$) score, female gender ($p=0.005$), and larger practice size ($p<0.001$) were positively associated with GP agreement that the patient did not have dementia. Subjective memory complaints ($p<0.001$) were more likely to result in a false positive diagnosis of dementia.

Conclusions: Timely recognition of dementia is advocated for optimal dementia management, but early recognition of a possible dementia syndrome needs to be balanced with awareness of the likelihood of false positives in detection. Although GPs correctly agree

with dimensions measured by the CAMCOG-R, improvements in sensitivity are required for

earlier detection of dementia.

Key words: cognition disorders, diagnosis, subjective memory complaint, primary care,

family practice, GP

Running title: GP detection of dementia and CAMCOG-R

Introduction

General practitioners (GPs) are frequently the first health professionals contacted by older

people and/or their family/carer(s) when symptoms of dementia cause concern (Speechly *et al.*, 2008). Dementia is a complex and variable condition (Hansen *et al.*, 2008; Palmer *et al.*, 2008a) which makes recognition of it particularly difficult in a low prevalence primary care setting (Pentzek *et al.*, 2009a). In the early stages, GPs need to make the often challenging distinction between normal aging, mild cognitive impairment, early dementia and/or cognitive impairment associated with depression, delirium or drugs (Pond and Brodaty, 2004). Timely detection and earlier recognition of dementia is desirable to optimise patient care and carer well-being (Phillips *et al.*, 2011). Conversely, a false positive diagnosis of dementia may result in medical over-investigation and overtreatment, lack of attention to manageable conditions such as depression, and unnecessary distress to patients and families. About 50% of patients over 65 with dementia are not diagnosed by GPs (Iliffe *et al.*, 2009b) for various reasons which include system characteristics and GP-related and patient-related factors (Koch and Iliffe, 2010). GPs' recognition of dementia appears to be associated with dementia severity (as measured by cognitive function tests and Instrumental Activities of Daily Living functioning) (Pentzek *et al.*, 2009b; Wind *et al.*, 1994) regardless of potential confounders such as depression, other illness, and recent events (van Hout *et al.*, 2007; Wind *et al.*, 1994). People with mild dementia and those who live alone are less likely to be diagnosed by their GP (Pentzek *et al.*, 2009b). Presumably those living alone lack informant reports which have been shown to be valuable in recognising patient's cognitive deficits (Jorm, 2004).

GPs also need to be cautious not to over-diagnose dementia. Factors associated with false-positive judgments of dementia include: higher age, patients' mention of memory complaints, GP-documented depression, lower education levels, and physical problems (e.g. mobility,

hearing) (Pentzek *et al.*, 2009a; Pentzek *et al.*, 2009b). Patient gender can also influence GP judgment of dementia, although findings are inconsistent and the effects are generally not marked (van Hout *et al.*, 2007; Wind *et al.*, 1994).

Patient mention of subjective memory complaints (SMC) can influence GPs' judgments, contributing to correct identification of dementia (Wind *et al.,* 1994) but also to over-diagnosis (Pentzek *et al.,* 2009b). Whilst patients' mention of SMC may place them at higher risk of being falsely judged as having dementia, patients with cognitive problems may not necessarily raise these as an issue with their GP (Waldorff *et al.*, 2008). People with early dementia can lack insight into their decline or may attribute symptoms such as memory problems to 'normal ageing'(Campbell *et al.*, 2008; Clare, 2003; Iliffe *et al.*, 2009a; Waldemar *et al.*, 2007). Controversy exists in the literature about the merit of patients' reports of SMC as a predictor or precursor of dementia (Coley *et al.*, 2008; Mitchell, 2008; Palmer *et al.*, 2008a; b; Pentzek *et al.*, 2009b; Reid and MacLullich, 2006), partly due to the range of non-standardised methods used to measure SMC, and their consistent association with depression and some personality traits (Reid and MacLullich, 2006). Although SMC appear to be predictive of future cognitive decline and/or dementia (Reid and MacLullich, 2006), they have modest diagnostic value and should not be relied upon for case-finding (Mitchell, 2008; Palmer *et al.*, 2008b). However, as a brief method of excluding healthy people in low prevalence settings, SMC seem to work reasonably well, that is, they exclude healthy elderly from those with dementia about 19 times out of every 20 non-cases (Mitchell, 2008).

It has been suggested recently that the process of dementia diagnosis should be split into four diagnostic steps (the trigger phase, disease oriented diagnosis, care-oriented diagnosis, carer assessment) followed by a monitoring phase (Buntix *et al.*, 2011). In this case the first stage of dementia diagnosis is not testing, but considering the possibility that dementia may

be emerging. GPs' suspicion and recognition of dementia rests on their clinical judgment based on their objective and subjective impressions from personal observation along with information from the patient and/or the patient's family/carer(s). Making the correct diagnosis is not always easy; many factors may interplay and can confound decisions

We aimed to use data from a larger study to examine predictors (in terms of GP and patient factors) of GP agreement with an objective assessment of cognitive impairment, the CAMCOG-R.

Methods

Participants

This study utilized baseline data collected from GPs and their patients as part of the "Ageing in General Practice" study (Pond *et al.*, 2012). Briefly, GPs and their patients were recruited from one rural and four metropolitan Australian sites: Sydney (NSW), Melbourne (Victoria), Adelaide (South Australia), Newcastle (NSW) and Bendigo (Victoria). GP practices within each study location were allocated a random approach order, and invited to participate in the study via phone. GPs who expressed an interest in participation were visited in order for the study to be explained. Basic demographics were collected at this visit from those GPs who agreed to participate.

Patients were recruited by a mail-out from each consenting GP to all community dwelling, English speaking patients aged 75 or older on their database. Patients with Parkinson's disease, multiple sclerosis, motor neuron disease, central nervous system inflammation, pre-existing psychotic symptoms, developmental disability, a history of substance abuse or a progressive malignancy were excluded.

GP Audit

GPs were sent a list of their participating patients by fax or email. They were asked to confirm each patient's eligibility for the study and to provide their clinical judgment in relation to each patient's dementia status using one of four options: No Dementia, Possible Dementia, Probable Dementia or Definite Dementia. GPs returned their completed audit forms to the local project officer via mail, fax or email. For any patients with a clinical judgment of dementia, GPs were asked whether the patient had completed a formal cognitive

function test or had been referred to a specialist. In Australia, specialist attendance is not reimbursed unless the patient has been referred by their general practitioner.

Patient Assessment

After receipt of the completed audit from the GP, patients were visited at home by a research nurse who collected demographic data from all consenting patients and assessed their cognitive function and depression status using the instruments described below. Patients were asked the following questions: "Do you have any complaints about your memory? Have you mentioned these to your GP?"

Revised Cambridge Cognitive Examination: Cognitive function was assessed using the CAMCOG-R subsection of the revised Cambridge Examination for Mental Disorders of the Elderly (CAMDEX-R) (Roth *et al.*, 1998). The CAMCOG-R consists of 68 questions covering 7 domains: Orientation (10 items), Language (17 items), Memory (13 items), Attention/Calculation (4 items), Praxis (8 items), Abstract Thinking (4 items) and Perception (3 items). The highest possible score is 104, with a cut-point of 79/80 indicative of dementia in lower scorers with 93% sensitivity and 87% specificity (Huppert *et al.*, 1996). The authors recognize that interpretation of CAMCOG-R scores around the cut-off is problematic. Although CAMCOG-R score is a significant predictor of dementia, it is insufficient for a clinical diagnosis of dementia (van Hout *et al.*, 2001). For the purposes of this study, a CAMCOG-R score of 79 or less was used as a sensitive and reasonably specific indicator of dementia.

Geriatric Depression Scale: Depression was measured using the short form 15-item Geriatric Depression Scale (GDS) developed by Sheikh and Yesavage (1986), that is widely used in primary health care settings for assessment of depression in the elderly (de Craen *et al.*,

2003). Scores above 5 on the short form of the GDS are considered to indicate depression, with 6 to 10 indicative of mild to moderate depression and 11 to 15 indicative of severe depression (Friedman *et al.*, 2005). A cut off-score of 5/6 was used in this study to distinguish those without depression from those participants with mild to moderate or severe depression.

GP Agreement with CAMCOG-R

The clinical judgment of the GP was classified as being in agreement with the CAMCOG-R if their dementia audit indicated (i) "No" for patients with a CAMCOG-R score greater than 79, or (ii) "Yes", "Possible" or "Probable" for their patients that scored less than 80 on the CAMCOG-R. The rationale for this categorization was that it was the most clinically relevant option: a judgment of "possible" dementia indicated that the patient had been flagged by the GP for surveillance (albeit informal) of worsening condition – the "trigger phase" of a dementia diagnosis (Buntix *et al.*, 2011) had occurred.

Statistical Analyses

Continuous data were summarized using the mean and standard deviation (SD). Categorical data were summarized as counts and percentage of agreement within each category. Consistency of agreement between the CAMCOG-R classification of dementia and GP suspicion or recognition of dementia was tested using Cohen's kappa. Logistic regression was used to examine the predictors of agreement of GPs' diagnoses and CAMCOG-R classifications separately in participants with and without CAMCOG-R dementia. The logistic regression model was fitted within a Generalized Estimating Equation (GEE) framework to adjust for the clustering of patients within GPs.

Ethics approval

Ethics approval was sought and granted initially from the Newcastle University Human Research Ethics Committee (Approval No. H-151-1205), and following this, from the appropriate Ethics Committees at each site.

Results

A total of 2,028 community dwelling patients aged 75 years or over were recruited via 169 GPs. The response rate was 6% for GPs and 19% for patients. Home assessment of cognitive and depression status (using CAMCOG-R and GDS) was completed for 2,024 patients. The dementia prevalence based on a CAMCOG-R cut score of 79/80 was 8.2%. Mild depression was present in 6.7% of patients (GDS score 6-10) and severe depression in 0.5% of patients (GDS score 11-15). Approximately 20% of patients indicated that they had expressed concerns about their memory to their GP (Table 1).

The participating GPs (n=165) had between 1 and 56 patients in the study. Four GPs were not included in the study, as none of their patients consented or completed the home assessment. A GP audit of dementia status was completed for 1,974 (97.5%) patients (Table 2). Of the 164 patients with CAMCOG-R dementia, 44.5% were identified by GPs as having dementia: 15.9% as "Definite", 9.1% as "Probable" and 19.5% as "Possible" dementia. GPs reported using a pencil and paper test of cognitive function for 63 of the 254 (24.8%) patients whom they judged to have dementia, and had referred 23 patients (9%) to a memory specialist.

GP Audit for Dementia

There was fair agreement between the CAMCOG-R and the clinical judgment of GPs in relation to the dementia status of patients, with GPs correctly identifying 73 (44.5%) of patients with and 1629 (90%) of patients without CAMCOG-R dementia (κ=0.276). Using our definition, (definite, probable and possible dementia) GPs also identified dementia in 181 people (11%) without CAMCOG-R dementia. This resulted in a positive predictive value (PPV) of 73/254 (28.7%) for definite, probable or possible dementia. That is, only 28.7% of

patients the GPs identified as having dementia were identified as such on the CAMCOG-R assessment. The sensitivity and specificity of GP judgment in relation to patient dementia status was 0.45 and 0.90 respectively.

The PPV of a GP judgment of "definite" dementia was 46%, and dropped to 31% and 21% for judgments of "probable" and "possible" respectively, substantiating lower levels of certainty amongst the GPs.

Patients with CAMCOG-R Dementia

Two patient dependent factors were most important for predicting agreement between the CAMCOG-R and GP judgment in patients with dementia: the CAMCOG-R score (p= 0.006) and patient mention of SMC to the GP ($p = 0.040$; Table 3). For dementia patients, the CAMCOG-R score was significantly lower for those with a GP clinical judgment of dementia (67.0 ± 12.2), compared to those judged to be cognitively intact by their GP (73.3 ± 7.5). Agreement between the GP and CAMCOG-R was approximately 20% greater in those patients with dementia that had mentioned a memory complaint to the GP (Table 3). Depression scores (GDS) were not significantly associated with GP diagnosis of dementia ($p=0.085$; Table 3).

Patients without CAMCOG-R Dementia

Patient factors. A higher CAMCOG-R score was significantly associated with GP agreement that the patient did not have dementia ($p<0.001$; Table 3). Two other patient factors were significant: gender, with the GP more likely to correctly identify the patient as not having dementia if they were female ($p=0.005$); and patient complaint about their memory ($p<0.001$). GPs were more likely to incorrectly identify the patient as having dementia if they complained about their memory. Thus, for patients without CAMCOG-R dementia, GP

agreement with the CAMCOG-R was greatest when the patient was female, did not mention a memory complaint, and scored higher on the CAMCOG-R (Table 3).

GP factors. Several GP factors were associated with correct identification of patients who did not have dementia. GPs from larger practices were more likely to agree with the CAMCOG-R in this group (p<0.001; Table 3). GPs who visited nursing home patients, were more likely to see dementia where it was not present (p=0.036) as were solo GPs (p<0.001). That is, GP agreement with the CAMCOG-R was greatest where GPs were from a larger practice and did not visit nursing home patients.

The characteristics of GP practices were examined by practice size (Table 4) in order to further examine the association of improved GP clinical judgment in larger practices. Larger practices were more likely to have a practice nurse ($p < 0.001$); their GPs in this study were younger ($p < 0.001$), and had spent less time as a practicing GP ($p < 0.001$). Almost 80% of solo GPs were male, but there was no significant relationship overall between GP gender and practice size (p=0.082).

Subjective memory complaints (SMC)

A total of 337 patients mentioned a concern about memory problems to their GP; 61 (18%) of these patients had CAMCOG-R dementia (Table 5). The sensitivity and specificity for diagnosing dementia on the basis of SMC expressed by the patient to their GP was 0.37 and 0.85 respectively. The positive and negative predictive values of SMC for CAMCOG-R dementia were 0.18 and 0.94 respectively. Calculation of a positive likelihood ratio from these values indicated that patients in this study who expressed SMC to their GP were 2.46 times more likely than the general population to have CAMCOG-R dementia.

Discussion

The current study investigated GP and patient factors that predicted GP agreement with an objective patient assessment for dementia (CAMCOG-R). GP agreement with the CAMCOG-R was fair, (κ= 0.276), due in large part to agreement between the GPs and the CAMCOG-R on absence of dementia. GPs correctly identified 90% of patients without CAMCOG-R dementia, though they were correct in less than half of their patients (44.5%) who had dementia, resulting in a relatively low PPV (28.7%). This is similar to the results of other studies of GPs or family physicians (Boustani *et al.*, 2003; Pentzek *et al.*, 2009a; Valcour *et al.*, 2000).

GPs were more likely to correctly identify dementia in those patients with a lower (i.e. poorer) score on the CAMCOG-R and more likely to correctly judge absence of dementia in those that scored well on the CAMCOG-R. Other studies have also reported that GPs' recognition of dementia was better in moderate to severe cases (Pentzek *et al.*, 2009a; Pentzek *et al.*, 2009b; van Hout *et al.*, 2007; Wind *et al.*, 1994). Relatively few GPs relied on specialist advice or the use of an objective assessment tool (e.g. Mini Mental State Exam) to assess patient cognition (used for 24.8% of patients), suggesting that their judgments were based on predominately subjective evidence. Impaired cognitive function and/or everyday functioning have previously been found to be associated with GP's subjective judgment on dementia (Wind *et al.*, 1994). Clearly, GPs are on the right track in agreeing with dimensions measured by the CAMCOG-R, though for earlier detection of dementia improvements in sensitivity are required.

Patients with SMC were 20% more likely to be diagnosed correctly as having dementia, though almost 20% less likely to be correctly identified as not having dementia. The positive and negative predictive values of SMC for CAMCOG-R dementia were 0.18 and 0.94

respectively, which is the same as previously reported for a low prevalence community sample (Mitchell, 2008). Memory complaints increased the likelihood of dementia by 2.46 times, from 8.2% to around 20%. Although SMC were reported by 56% of people with CAMCOG-R dementia in this study, only 37% had mentioned it to their GP. These GPs clearly responded to SMC cues from their patients to identify dementia, however they also overrated the presence of dementia more frequently in patients with SMC, a finding in accord with previous research (Pentzek *et al.*, 2009b; Wind *et al.*, 1994). Thus, while helpful in making the diagnosis for those patients who have dementia, SMC also contribute to over-diagnosis in those who do not. In at least some cases though, SMC may be predictive of future cognitive decline (Geerlings *et al.*, 1999; Reid and MacLullich, 2006).

In a community setting, even when patients report SMC to their GP, there is only a 20% chance that dementia is present, but these people may have up to a threefold increased risk of developing Alzheimer's disease in the future (Geerlings *et al.*, 1999). A population-based study in Sweden reported that only 9-15% of participants with memory or multiple impairments did not have subjective memory complaints (Palmer *et al.*, 2008b). It may be that GPs are sensitive to this possibility. Although SMC alone have modest diagnostic value for dementia, in combination with an objective assessment they may offer advantage over either method alone, and provide a mechanism for improvement of GP diagnosis of dementia.

In addition to lower CAMCOG-R score and SMC, the other patient characteristic related to a false positive diagnosis of dementia was gender. There were fewer false positive GP judgments of dementia for female patients (9%) compared to male patients (11%). This small but significant effect may be related to differences in the length and type of GP consultation between female and male patients. GPs spend longer on average with female patients (Britt *et al.*, 2005), and spend a higher percentage of time on physical examination, screening, patient questions, and emotional counseling, whereas visits by men involve a

higher percentage of time spent on procedures and health behavior counseling (Tabenkin *et al.*, 2004). The dynamics of the GP consultation with female patients may be more conducive to a correct ruling-out of dementia.

GP factors that were associated with an increased false positive diagnosis of dementia were practice size, and regular visits to nursing home patients. GPs from larger practices were more likely to correctly rule-out dementia. Interestingly, Connolly *et al* (2011) reported lower rates of diagnosis in practices with one GP as opposed to several. They considered one of the reasons for this may have been lack of time for the solo GP. Lack of time may also be a factor in the over-diagnosis of dementia by solo GPs in our study, as larger practices were more likely to have a practice nurse who may assist with dementia assessments. Larger practices were also more likely to employ younger GPs who may have received more dementia education. In the Australian context, GPs in solo practice and those who do nursing home visits tend to be older (Charles *et al.*, 2006; Gadzhanova *et al.*, 2007).

GPs that did nursing home visits were also more likely to over-diagnose dementia, an interesting finding that is difficult to explain. Possibly their exposure to dementia in nursing homes may have influenced their perception of the incidence of dementia in the aged population. That is, GPs with regular exposure to the nursing home environment may perceive dementia in the aged to be a common and likely occurrence, as dementia is the most common problem managed in Australian nursing homes by GPs – 33 times the usual management rate in everyday practice (O'Halloran *et al.*, 2007).

Limitations of this study include the self-selected nature of the GP and patient populations and the use of a single cut point on the CAMCOG-R as the only benchmark for a correct diagnosis of dementia. Although the CAMCOG-R is a significant predictor for the clinical diagnosis of dementia (van Hout *et al.*, 2001), most cases require additional clinical judgment for a definitive diagnosis. GPs may have been sensitive to patient presentation that suggested

deficits relating to factors such as activities of daily living and the mention of memory problems which may not be reflected in a cognitive function test, as only 28.7% of patients with a GP judgment of dementia had CAMCOG-R dementia. This may have inflated the false positive diagnoses found in this study.

While timely recognition of dementia is advocated for optimal dementia management, at present the early recognition of a possible dementia syndrome needs to be balanced with awareness of the likelihood of false positives in detection. Clearly, GPs are on the right track in agreeing with dimensions measured by the CAMCOG-R, though for earlier detection of dementia improvements in sensitivity are required. Enquiring about memory issues and monitoring patients voicing SMC is a consideration, though using this enquiry for diagnosis raises the risk of increasing the false positive rate. Screening tests have been advocated but are also associated with a poor positive predictive value (Brodaty *et al.*, 2006). A more complex approach is needed, perhaps including the use of pretest probabilities such as that provided by memory complaints.

Conflict of Interest Declaration

CDP: on advisory boards for dementia for a number of companies, including in the last 3 years Pfizer, Lundbeck and Nutricia, all of which have been paid positions; received sponsorship to attend conferences from Pfizer; been a speaker for the Dementia Study Training Centres and for Alzheimers Australia and received honoraria for these activities; received grants fron the NHMRC and Department of Health and Ageing in relation to dementia.has sat on advisory boards for Pfizer, Novartis, Janssen and Lundbeck.

HB: received grants from pharmaceutical companies to be a participating center in multicenter international drug trials for Alzheimer's disease (currently Sanofi, Servier, Roche); has received honoraria from Pfizer, Janssen, Lundbeck, Novartis for giving presentations or chairing meetings; has been a sponsored speaker by Pfizer, Janssen, and Novartis; and has been a consultant and/or sat on advisory boards for Pfizer, Janssen, Lundbeck, Novartis, Merck. Nutricia and Lilly.

All other authors have no competing interests to declare.

Description of Authors' Roles

Dimity Pond conceived and developed the study and contributed to writing the paper. Karen Mate assisted with design of the study, supervised the data collection and contributed to writing the paper. Jill Phillips assisted with writing the paper. Natasha Weaver did the statistical analyses. Parker Magin, Nigel Stocks and Henry Brodaty assisted in study design and writing of the paper.

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References

Boustani, M., Peterson, B., Hanson, L., Harris, R. and Lohr, K. (2003). Screening for Dementia in Primary Care: A Summary of the Evidence for the U.S. Preventive Services Task Force. *Annals of Internal Medicine*, 138, 927-937.

Britt, H., Valenti, L. and Miller, G. (2005). Determinants of consultation lenght in Australian general practice. *Medical Journal of Australia*, 183, 68-71.

Brodaty, H., Low, L. F., Gibson, L. and Burns, K. (2006). What is the best dementia screening instrument for general practitioners to use? *American Journal of Geriatric Psychiatry*, 14, 391-400.

Buntix, F., De Lepeleire, J., Paquay, L., Iliffe, S. and Schoenmakers, B. (2011). Diagnosing dementia: No easy job. *BMC Family Practice*, 12, 60-64.

Campbell, K. H., *et al.* (2008). Dementia, diagnostic disclosure, and self-reported health status. *Journal of the American Geriatrics Society*, 56, 296-300.

Charles, J., Britt, H., Valenti, L., Charles, J., Britt, H. and Valenti, L. (2006). The independent effect of age of general practitioner on clinical practice. *Medical Journal of Australia*, 185, 105-109.

Clare, L. (2003). Managing threats to self: awareness in early stage Alzheimer's disease. *Social Science and Medicine*, 57, 1017-1029.

Coley, N., Ousset, P. J., Andrieu, S., Matheix Fortunet, H. and Vellas, B. (2008). Memory complaints to the general practitioner: data from the GuidAge study. *Journal of Nutrition, Health and Aging*, 12, 66S-72S.

Connolly, A., Gaehl, E., Martin, H., Morris, J. and Purandare, N. (2011). Underdiagnosis of dementia in primary care: Variations in the observed prevalence and comparisons to the expected prevalence. *Aging and Mental Health*, 15, 978-984.

de Craen, A., Heeren, T. and Gussekloo, J. (2003). Accuracy of the 15-item Geriatric Scale (GDS-15) in a community sample of the oldest old. *International Journal of Geriatric Psychiatry*, 18, 63-66.

Friedman, B., Heisel, M. and Delavan, R. (2005). Psychometric Properties of the 15-Item Geriatric Depression Scale in Functionally Impaired, Cognitively Intact, Community-Dwelling Elderly Primary Care Patients. *Journal of the American Geriatrics Society*, 53, 1570-1576.

Gadzhanova, S., Reed, R., Gadzhanova, S. and Reed, R. (2007). Medical services provided by general practitioners in residential aged-care facilities in Australia. *Medical Journal of Australia*, 187, 92-94.

Geerlings, M. I., Jonker, C., Bouter, L. M., Ader, H. J. and Schmand, B. (1999). Association between memory complaints and incident Alzheimer's Disease in elderly people with normal baseline cognition. *American Journal of Psychiatry*, 156, 531-537.

Hansen, E. C., Hughes, C., Routley, G. and Robinson, A. L. (2008). General practitioners' experiences and understandings of diagnosing dementia: Factors impacting on early diagnosis. *Social Science and Medicine*, 67, 1776-1783.

Huppert, F. A., *et al.* (1996). Psychometric properties of the CAMCOG and its efficacy in the diagnosis of dementia. *Aging, Neuropsychology and Cognition*, 3, 201-214.

Iliffe, S., Jain, P. and Wilcock, J. (2009a). Recognition of and response to dementia in primary care:part 1. *InnovAit*, 2, 230-236.

Iliffe, S., *et al.* (2009b). Primary care and dementia: 1. diagnosis, screening and disclosure. *International Journal of Geriatric Psychiatry*, 24, 895-901.

Jorm, A. (2004). The informant questionnaire on cognitive decline in the elderly (IQCODE): a review. *International Psychogeriatrics*, 16, 275-293.

Koch, T. and Iliffe, S. (2010). Rapid appraisal of barriers to the diagnosis and management of patients with dementia in primary care: a systematic review. *BMC Family Practice*, 11.

Mitchell, A. J. (2008). The clinical significance of subjective memory complaints in the diagnosis of mild cognitive impairment and dementia: a meta-analysis. *International Journal of Geriatric Psychiatry*, 23, 1191-1202.

O'Halloran, J., Britt, H., Valenti, L., O'Halloran, J., Britt, H. and Valenti, L. (2007). General practitioner consultations at residential aged-care facilities. *Medical Journal of Australia*, 187, 88-91.

Palmer, K., *et al.* (2008a). Early symptoms and signs of cognitive deficits might not always be detectable in persons who develop Alzheimer's disease. *International Psychogeriatrics*, 20, 252-258.

Palmer, K., et al. (2008b). Mild cognitive impairment in the general population: occurrence and progression to Alzheimer disease. *American Journal of Geriatric Psychiatry*, 16, 603-611.

Pentzek, M., et al. (2009a). General practitioners' judgment of their elderly patients' cognitive status. *Journal of General Internal Medicine*, 24, 1314-1317.

Pentzek, M., et al. (2009b). Apart from nihilism and stigma: what influences general practitioners' accuracy in identifying incident dementia? *American Journal of Geriatric Psychiatry*, 17, 965-975.

Phillips, J., Pond, D. and Goode, S. (2011). Timely diagnosis of dementia: Can we do better. Sydney: Alzheimer's Australia.

Pond, C., et al. (2012). Ageing in general practice (AGP) trial: a cluster randomised trial to examine the effectiveness of peer education on GP diagnostic assessment and management of dementia. *BMC Family Practice*, 13, 12.

Pond, D. and Brodaty, H. (2004). Diagnosis and management of dementia in general practice. *Australian Family Physician*, 33, 789-793.

Reid, L. M. and MacLullich, A. M. (2006). Subjective Memory Complaints and Cognitive Impairment in Older People. *Dementia and Geriatric Cognitive Disorders*, 22, 471-485.

Roth, M., Huppert, F. A., Mountjoy, C. Q. and Tym, E. (1998). *CAMDEX-R: The Cambridge examination for mental disorders of the elderly - revised*. Cambridge: Cambridge University Press.

Sheikh J. I. and Yesavage, J. A. (1986). Geriatric Depression Scale (GDS): Recent evidence and development of a shorter version. In T. L. Brink (Ed.) Clinical Gerontology: A Guide to Assessment (pp. 165-173) New York: Haworth Press.

Speechly, C. M., Bridges-Webb, C. and Passmore, E. (2008). The pathway to dementia diagnosis. *Medical Journal of Australia*, 189, 487-489.

Tabenkin, H., Goodwin, M., Zyzanski, S., Stange, K. and Medalie, J. (2004). Gender differences in time spent during direct observation of doctor-patient encounters. *Journal of Women's Health*, 13, 341-349.

Valcour, V. G., Masaki, K. H., Curb, J. D. and Blanchette, P. L. (2000). The detection of dementia in the primary care setting. *Archives of Internal Medicine*, 160, 2964-2968.

van Hout, H., et al. (2001). CAMDEX, can it be more efficient? Observational study on the contribution of four screening measures to the diagnosis of dementia by a memory clinic team. *International Journal of Geriatric Psychiatry*, 16, 64-69.

van Hout, H. P. J., Vernooij-Dassen, M. J. and Stalman, W. A. B. (2007). Diagnosing dementia with confidence by GPs. *Family Practice*, 24, 616-621.

Waldemar, G., et al. (2007). Access to diagnostic evaluation and treatment for dementia in Europe. *International Journal of Geriatric Psychiatry*, 22, 47-54.

Waldorff, F. B., Rishoj, S. and Waldemar, G. (2008). If you don't ask (about memory), they probably won't tell. *The Journal of Family Practice*, 57, 41-44.

Wind, A. W., van Staveren, G., Schellevis, F. G., Jonker, C. and et al. (1994). The validity of the judgement of general practitioners on dementia. *International Journal of Geriatric Psychiatry*, 9, 543-549.

Table 1. Characteristics of patient and GP participants

Patients (n=2,024)	
Age (years) [mean ± sd]	81.3 ± 4.2
Gender: Female	54.5%
Marital Status: Married/de facto	50.8%
Divorced/Separated	7.0%
Single/Never married	4.8%
Widow	37.4%
IRSAD[‡]	7.0 ± 2.5
Mentioned memory problems to GP	19.9%
CAMCOG performance	90.2 ± 8.0
Cognitively impaired on CAMCOG-R	8.2%
GDS Score	2.1 ± 2.1
Depression (GDS): Mild to Moderate	6.7%
Severe	0.5%
General Practitioners (n=165)	
Age (years)	50.8 ± 8.9
Time as GP (years)	21.9 ± 9.1
Gender: Female	41.3%
No of patients in study	11.3 ± 10.2
Practice Size: Solo	16.9%
2-4 GPs	31.3%
≥ 5 GPs	45.6%
Practice has a Practice Nurse: Yes	56.3%
GP Visits Nursing Home Patients: Yes	75.6%
GP Routinely Performs 75+ Check: Yes	68.8%
Additional Education [†]	6.7%

[†] Extra qualifications in mental health or geriatrics
[‡] Index of Relative Socioeconomic Advantage and Disadvantage (scale 1-10, with 10 being most advantaged).

Table 2. GP judgment of dementia status compared to CAMCOG-R result

GP Judgment of Dementia Status	CAMCOG-R Result[a]	
	Dementia n =164 (%)	No Dementia n=1810 (%)
Definite	26 (15.9)	31 (1.7)
Probable	15 (9.1)	33 (1.8)
Possible	32 (19.5)	117 (6.5)
No	91 (55.5)	1629 (90)

[a] Patients who scored < 80 on the CAMCOG-R were classified as having dementia.

Table 3. Predictors of agreement between GP clinical judgment and the CAMCOG-R assessment of dementia

	Dementia (CAMCOG-R score < 80)			No Dementia CAMCOG-R score ≥ 80)		
	GP Judgment		P-value	GP Judgment		P-value
	Disagree n=91 (%)	Agree n=73 (%)	(GEE)	Disagree n=181 (%)	Agree n=1629 (%)	(GEE)
Patient Factors						
Gender			0.372			0.005
Male	41 (55)	34 (45)		91 (11)	724 (89)	
Female	50 (56)	39 (44)		90 (9)	903 (91)	
Age	83 ± 5.0	84.1 ± 4.6	0.383	82.4 ± 4.8	80.9 ± 4.0	0.123
Marital Status			0.183			0.228
Married/defacto	39 (46)	45 (54)		87 (9)	829 (91)	
Other	51 (65)	28 (35)		10 (8)	120 (92)	
Mention Memory to GP			0.040	67 (25)	201 (75)	<0.001
Yes	25 (42)	34 (58)				
No	65 (63)	39 (38)		114 (7)	1424 (93)	
CAMCOG-R	73.3 ± 7.5	67.0 ± 12.2	0.006	89.6 ± 5.3	92.1 ± 4.8	<0.001
GDS	3.6 ± 3.2	3.1 ± 2.5	0.085	2.5 ± 2.2	1.9 ± 1.9	0.201
GP Factors						
Gender			0.133			0.100
Male	58 (60)	39 (40)		89 (8)	1023 (92)	
Female	27 (49)	28 (51)		80 (13)	542 (87)	
Age	53.4 ± 8.2	50.8 ± 7.5	0.120	51.0 ± 7.0	52.0 ± 8.0	0.142
Practice Size			0.391	55 (15)	301 (85)	<0.001
Solo	17 (74)	6 (26)				
2-4 GPs	32 (52)	29 (48)		64 (11)	500 (89)	
More than 5 GPs	35 (54)	30 (46)		39 (5)	712 (95)	
Nursing Home Patients			0.598	141 (10)	1248 (90)	0.036
Yes	70 (55)	58 (45)				
No	14 (67)	7 (33)		17 (6)	265 (94)	
Performs 75+ Check			0.854	106 (9)	1141 (91)	0.180
Yes	60 (59)	42 (41)				
No	24 (51)	23 (49)		52 (13)	360 (87)	

Table 4. Characteristics of general practices based on practice size

Characteristic	Practice Size (Number of GPs)			P-values
	1	2-4	≥ 5	
	n=27 (%)	n=50 (%)	n=75 (%)	
GP Age	58.3 (6.9)	50.2 (8.1)	48.6 (8.7)	<0.001
GP Gender				
Male	21 (78)	27 (54)	41 (55)	0.0820
Female	6 (22)	23 (46)	34 (45)	
GP Time	30.1 (7.2)	21.1 (8.3)	19.4 (8.6)	<0.001
Practice Nurse				
Yes	4(15)	27 (54)	61 (81)	<0.001
No	23 (85)	23 (46)	14 (19)	
Practice does 75+ check				
Yes	18 (69)	33 (66)	61 (81)	0.1299
No	8 (31)	17 (34)	14 (19)	
Visits nursing home patients				
Yes	23 (85)	35 (70)	64 (85)	0.0840
No	4 (15)	15 (30)	11 (15)	

Table 5. Relationship between subjective memory complaint and CAMCOG-R result

Subjective Memory Complaint	CAMCOG-R Result		Total
	Impaired n=166 (%)	Not Impaired n=1857 (%)	
Yes	93 (11)	747 (89)	840
No	73 (6)	1108 (94)	1181
Mentioned to GP[†]			
Yes	61 (18)	276 (82)	337
No	105 (6)	1581 (94)	1686

[†]Based on participant report that they had discussed memory complaint with their GP