

THE MINI-COG: RECEIVER OPERATING CHARACTERISTICS WITH EXPERT AND NAÏVE RATERS

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ABSTRACT

Background. As elderly populations grow, dementia detection in the community is increasingly needed. Existing screens are largely unused because of time and training requirements. We developed the Mini-Cog, a brief dementia screen with high sensitivity, specificity, and acceptability. Here we describe the development of its scoring algorithm, its receiver operating characteristics (ROC), and the generalizability of its clock drawing scoring system.

Sample and methods. A total of 249 multi-lingual older adults were examined. Scores on the **three-item recall** task and the **clock drawing task (CDT-CERAD version)** were combined to create an optimal algorithm. Receiver operating characteristics for seven alternatives were compared with those of the MMSE and the CASI using expert raters. To assess the CDT scoring generalizability, 20 naïve raters, without explicit instructions or prior CDT exposure, scored 80 randomly selected clocks as 'normal' or 'abnormal' (20 from each of four CERAD categories).

Results. An algorithm maximizing sensitivity and correct diagnosis was defined. Its ROC compared favorably with those of the MMSE and CASI. CDT concordance between naïve and trained raters was >98% for normal, moderately and severely impaired clocks, but lower (60%) for mildly impaired clocks. Recalculation of the Mini-Cog's performance, assuming that naïve raters would score all mildly impaired CDTs in the full sample as normal, retained high sensitivity (97%) and specificity (95%).

Conclusion. The Mini-Cog algorithm performs well with simple clock scoring techniques. The results suggest that the Mini-Cog may be used successfully by relatively untrained raters as a first-stage dementia screen. Further research is needed to characterize the Mini-Cog's utility when population dementia prevalences are low. Copyright © 2001 John Wiley & Sons, Ltd.

KEY WORDS—dementia screening; MMSE; clock drawing; three-item recall; Cognitive Abilities Screening Instrument; naïve raters; language

INTRODUCTION

In a recent survey of 368 general practitioners, 82% believed that dementia screening of elderly persons is worthwhile, but only 24% routinely screened their own patients (Bush *et al.*, 1997). The major barriers were lack of time (85%), fear of offending patients (58%) and inadequacies in available tests (22%). When a formal cognitive screen was used, it was usually the MMSE (Bush *et al.*, 1997); if a

short, simple screen such as clock drawing could be shown to be effective, 93% would use it.

Persuading doctors to adopt routine dementia screening thus appears to require a more efficient approach. The ideal instrument would be very brief, simple, sensitive, acceptable to older persons, and uninfluenced by low education and language barriers that weaken the utility of most available screens. The Mini-Cog is a new instrument that combines a simple memory test with clock drawing (CDT) and appears to fulfill these criteria. When used by expert raters, the Mini-Cog attained 99% sensitivity and 93% specificity in a split sample containing 50% demented persons, and required only 3 minutes to administer (Borson *et al.*, 2000). However, it is unclear whether novice raters unfamiliar with cognitive testing will accurately

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give the Mini-Cog in 'real world' settings. Its structure is simple enough to be used by community volunteers and health care personnel, provided its most vulnerable element, the CDT, can be scored correctly.

Many CDT dementia screening systems have been proposed, with varying degrees of complexity (Watson *et al.*, 1993; Shulman *et al.*, 1986; Dastoor *et al.*, 1991; Tuokko *et al.*, 1992; Death *et al.*, 1993). While all studies report the sensitivity and specificity of their proposed systems, cost-benefit, measured in ease of use, is rarely examined. If a scoring system slightly improves sensitivity or specificity but increases costs in terms of complexity, interpretation and training, then many CDT advantages are lost. As analog clock form and function are highly learned in most modern cultures, we believe that understanding how to judge CDTs as 'normal' and 'abnormal' is intuitively and widely understood. Therefore, if naïve raters gave their best estimates without an explicit scoring system, their judgments should closely resemble expert decisions. If true, this could greatly facilitate CDT use as in community-based dementia screening efforts.

Similar reasoning has been applied to the MMSE, and some studies show item subsets to perform as well as the complete scale. In one study, the four MMSE items that best discriminated normal controls from mild AD patients were two time orientation items (day, date) and two of the three conventional word recall items ('apple' and 'penny'). Dementia discrimination using these items alone was comparable to that of the full MMSE (Fillenbaum *et al.*, 1994). The MMSE three-item recall task was highly predictive of total score: 97% of subjects recalling 0/3 words had MMSE scores below the conventional cut-off point for screening (≤ 23), while 86% of those recalling 1-3 words scored ≥ 24 (Braekhus *et al.*, 1992). In five different studies designed to distill the essential MMSE items (Magaziner *et al.*, 1987; Roca, 1987; Klein *et al.*, 1985; Braekhus *et al.*, 1992; Fillenbaum *et al.*, 1994), only three-item recall was consistently implicated in both predicting total score and distinguishing normal from mildly demented subjects. Additionally, we have previously reported (Borson *et al.*, 2000) that three-item recall by itself was a sensitive (91%) and specific (97%) dementia discriminator. Accordingly, we combined this short learning task with the CDT to yield a dementia screen (the Mini-Cog) more sensitive than three-item recall

alone and relatively uninfluenced by education or language (Borson *et al.*, 2000). In this paper, we report the receiver operating characteristics (ROC) of different Mini-Cog scoring algorithms, and compare them with the MMSE and CASI. We also examine the accuracy of CDT scoring by untrained raters without scoring instruction, and whether Mini-Cog subject classification would be compromised by naïve rater scoring errors.

METHODS

Subjects

Full details of sample selection, subject characteristics, and diagnostic evaluations are provided elsewhere (Borson *et al.*, 1999, 2000). Briefly, 249 community dwelling older adults (173 women, 76 men) reflecting the five major ethnic groups in the United States completed the CDT, MMSE, and CASI during initial evaluation. All subjects or their proxies gave written informed consent using a multiple-language protocol approved by the University of Washington IRB.

Subjects were initially classified as 'probably demented' ($n=129$) or 'probably not demented' ($n=120$) based on an informant's history of cognitive decline (CERAD expanded history) and current functioning (Clinical Dementia Rating) (Hughes *et al.*, 1982), an approach analogous to that of Jorm *et al.* (1991). The classification of all subjects was subsequently confirmed using formal diagnostic criteria (CERAD, DSM-IV: American Psychiatric Association, 1994; and NINCDS-ADRDA: McKhann *et al.*, 1984), and demented subjects were further diagnosed as having probable or possible AD, another dementia, or no dementia. Subjects with uncertain/very mild cognitive impairment (CDR=0.5) were excluded. *Post hoc* dementia diagnoses were probable AD in 92 (71%), possible AD (mixed states, usually vascular brain disease in addition to AD) in 16 (12%), vascular dementia in 13 (10%), and other dementias in 8 (6%).

The CDT was scored using CERAD templates (Borson *et al.*, 1999). Subjects were instructed to draw a large circle, fill in the numbers on a clock face, and set the hands at 8:20, with no time limit. Excellent inter-rater CDT reliability was obtained (intra-class correlation=0.97) by two independent raters blind to each other's scores and to any subject information using the four-point CERAD scale (0, 1, 2, 3: 0=normal; 3=severe impairment).

While the three-item recall task can be given in both cued and uncued fashion, research suggests that uncued recall is sensitive to dementia in AD patients, and that cueing does not improve scores or dementia detection (Yuspeh *et al.*, 1998). Accordingly, we used uncued recall in this data analysis. One point was given for each word correctly recalled.

Clock scoring by naïve raters

We randomly sampled 20 CDTs from each of the four CERAD performance levels (0=normal, 1=mild, 2=moderate, 3=severe) for which exact agreement was reached by the two expert raters. The 80 CDTs were scrambled in random sequence. Twenty volunteers with no prior cognitive testing experience were recruited from hospital clinics, social service agencies, and administrative services. Naïve raters consisted of 9 (45%) administrative/clerical workers, 7 (35%) clinical/medical employees, and 4 (20%) community case managers. These raters averaged 39+10 years of age, with 16+2 years of education. All scored each CDT as normal or abnormal without hints or templates to guide them.

Data analyses

A series of Mini-Cog scoring algorithms were tested using various combinations of three-item recall and the CDT, to construct an ROC curve that was superimposed upon ROC curves generated for the MMSE and CASI using all possible

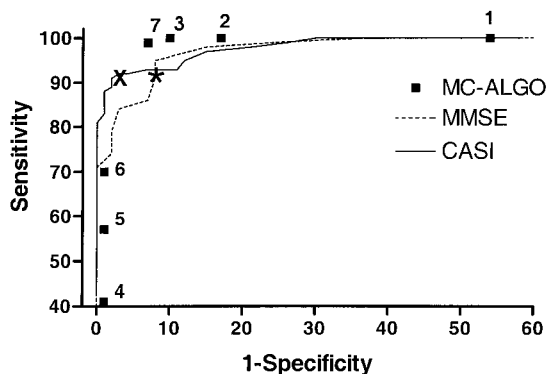


Fig. 1. ROC curves for MMSE and CASI, with M-C algorithms superimposed. 1-7 corresponds to Table 2 algorithms. X=conventional cut-off point for CASI (80); *conventional cut-off for MMSE (24)

Table 1. Correct classification of subjects by three-item recall and CDT scores (frequency and % shown)

CERAD CDT score	Number of words correctly recalled			
	0	1	2	3
0 ND (freq.)	3 (10)	9 (100)	34 (100)	65 (100)
DEM (freq.)	27 (90)	0 (0)	0 (0)	0 (0)
1 ND (freq.)	0 (0)	2 (100)	0 (0)	3 (100)
DEM (freq.)	16 (100)	0 (0)	3 (100)	0 (0)
2 ND (freq.)	0 (0)	0 (0)	1 (100)	1 (100)
DEM (freq.)	21 (100)	1 (100)	0 (0)	0 (0)
3 ND (freq.)	1 (2)	0 (0)	1 (25)	0 (0)
DEM (freq.)	53 (98)	3 (100)	4 (75)	1 (100)

CDT: 0=normal, 1=abnormal; ND=probably not demented (psychiatric judgement); D=probably demented (psychiatric judgement). Bold numbers indicate clearly demented (by Mini-Cog algorithm 7). Italic numbers indicate demented by Mini-Cog (would be classified as non-demented by naïve CDT raters). The remainder are clearly non-demented (by Mini-Cog).

cut-off points. In the formulation of Mini-Cog algorithms, preference was given to robust and simple combinations. The best available three-item/CDT combination was assessed against the conventional MMSE and CASI conventional cut-off points (MMSE=23/24 of 30 possible points (Folstein *et al.*, 1975); CASI=80/81 of 100 possible points (Teng *et al.*, 1994). Judgements of CDTs by naïve raters were compared with those of expert raters. Because naïve raters were simply asked to judge the CDTs as 'normal' or 'abnormal' and experts used the entire CERAD scale (0-3) we defined agreement as 'normal'=CERAD 0 and 'abnormal'=CERAD 1-3.

RESULTS

Algorithm construction

Cross-tabulations of three-item recall scores and CERAD CDTs were examined as the basis for constructing a classification algorithm when both tests were combined into a single instrument (Table 1). Various decision algorithms were constructed (see Table 2). Algorithm numbers correspond to those represented on the ROC curve as single points (Fig. 1) for comparison with the MMSE and CASI (represented as continuous curves). Results show that at least one recall/CDT algorithm had superior sensitivity relative to the CASI and MMSE. The conventional cut-off points for the

Table 2. Alternative Mini-Cog algorithms

Algorithm	Judged non-demented	Judged demented
1	Recall=3 and CDT=0	Recall=0-2 or CDT=1-3
2	Recall=2-3 and CDT=0	Recall=0-1 or CDT=1-3
3	Recall=1-3 and CDT=0	Recall=0 or CDT=1-3
4	Recall=1-3 or CDT=0-2	Recall=0 and CDT=3
5	Recall=1-3 or CDT=0-1	Recall=0-1 and CDT=2-3
6	Recall=1-3 or CDT=0	Recall=0 and CDT=1-3
7 Mini-Cog	Recall=3 or (Recall=1-2 and CDT=0)	Recall=0 or (Recall=1-2 and CDT=1-3)

MMSE and CASI corresponded relatively well with their points of optimal function (the best balance of sensitivity and specificity) in this sample.

When the performance of algorithm 7 (Table 3, henceforth referred to as the Mini-Cog) was compared with the other dementia screens (three-item alone, CDT alone, CASI, and MMSE), it performed best in sensitivity (99%) and diagnostic value (96%) and had acceptable specificity (93%), though it was less efficient in classifying non-demented subjects than the CASI (96%). The MMSE was the least sensitive (91%) and specific (92%) (Borson *et al.*, 2000). When separate logistic regression analyses predicting correct diagnosis and controlling for the effects of education and non-native language (English/non-English) were performed the results were striking. The Mini-Cog was most predictive ($\chi^2=273$), followed by the CASI ($\chi^2=233$) and the MMSE ($\chi^2=201$; see Borson *et al.*, 2000, for detailed description).

weaker for category 1 (60%) (Table 3). The 40% of naïve ratings discordant for mildly impaired CDTs, as judged by research raters, were uniformly rated as normal. Overall concordance between naïve and expert raters was 89%.

We evaluated the projected Mini-Cog performance as if it had been administered by naïve raters to our entire subject sample. We made the following assumptions: that the naïve raters properly administered the three-item recall task, that subjects received appropriate CDT instructions, that all mildly impaired CDTs were scored normal (rather than the 40% found here), and all other CDTs were scored identically to research ratings (rather than the 98–99% found here). Under these circumstances, Mini-Cog sensitivity would decrease from 99 to 97%, and specificity would increase from 93 to 95%, both well within effective screening ranges. The percentage of subjects with mildly impaired CDTs in this sample was small (10%), and 67% of these recalled 0/3 words.

CDT classification by naïve subjects

CDT judgments by naïve raters (normal or abnormal) were compared with expert judgments using CERAD categories (0, 1, 2, 3). Concordance between naïve and expert raters was extremely high for CERAD categories 0, 2 and 3 (>98%) and

DISCUSSION

We have previously reported that the combination of the three-item recall task with the CDT results in superior dementia prediction relative to either of its parts and to the CASI and MMSE in a very heterogeneous community sample (Borson *et al.*, 1999). Inspection of the ROC curves shown here further explicates those findings. First, the conventional cut-off points for the MMSE (23/24) and CASI (80/81) worked as expected in heterogeneous samples of older adults, producing the best sensitivity/specificity combination attainable with these instruments in this sample. Examining our algorithms (Table 2 and Fig. 1) we see that numbers 1–3 tend to maximize sensitivity, while 4–6 maximize specificity and 7 (the Mini-Cog) is a

Table 3. Naïve vs expert CDT ratings

Naïve raters	Expert raters CERAD categories			
	0	1	2	3
Normal	392* (98)†	160 (40)	8 (2)	1 (0.2)
Abnormal	8 (2)	240 (60)	392 (98)	399 (99.8)

*Absolute frequency based on N=400 (20 naïve raters×20 CDTs within each CDT category).

†Percentages.

hybrid of the two approaches. While there might be circumstances in which one attribute (sensitivity or specificity) might be more desirable than the other, we chose an approach that yielded the best overall percentage correct diagnosis, and high sensitivity with very good specificity (see Borson *et al.*, 2000, for further details). The optimal Mini-Cog algorithm was simple and robust, and used the simplest possible CDT scoring system: CERAD 0=normal, CERAD 1–3=abnormal.

Why does this simple approach work? The majority of factor analytic studies of the MMSE found two-factor solutions (Hill and Backman, 1995) including memory/recent learning as one factor; the second factor varying but usually including the visual-spatial MMSE components (Hill and Backman, 1995; Braekhus *et al.*, 1992). Three-item recall is a very simple task primarily testing short-term memory. We previously demonstrated its importance by showing that three-item recall alone yielded virtually the same correct diagnosis rate as the much longer CASI (Borson *et al.*, 2000). In contrast, algorithm calculations suggest that while the CDT alone is not as powerful a dementia screen as three-item recall, it improves the sensitivity and overall correct diagnosis achieved by the Mini-Cog. Accurate clock drawing requires many faculties impaired in dementia, including long-term memory, visuo-spatial representation, global attention, and executive functions. Combining these two elements seems to capture the essence of dementia for screening purposes.

Virtually all CDT studies show that greater impairment in clock drawing is associated with more severe dementia and lower MMSE scores (Death *et al.*, 1993; Brodaty and Moore, 1997; Juby, 1999). However devised, the cut-offs for these systems are usually set so that a small number of errors (10–25%) is consistent with non-demented status. For example, in the 10-point Shulman scale (Shulman *et al.*, 1986), a score of 8 was consistent with clinically non-demented status, while a score of ≤ 7 indicated possible dementia. With an alternative four-point system tested by Lam in a Chinese sample (Lam *et al.*, 1998), dementia screening cut-offs were set at 3 of 4 possible points. **The four-category CERAD system we used, while simpler than some alternative systems, uses a comparable cut-off point (0=normal, 1–3=demented).**

The CDT judgments by naïve raters were strikingly similar to those of expert raters for CERAD categories 0 (normal), 2 (moderate) and 3

(severe). The lower sensitivity of naïve than expert raters to mild CDT impairment (CERAD category 1) appeared unlikely to compromise Mini-Cog performance, even if failure to detect mild impairment was assumed to be complete. Using the full Mini-Cog algorithm, most would be classified as demented, even if all their CDTs were scored as 'normal', because the majority of subjects with mild CDT impairment (67%) scored poorly on three-item recall (0 of 3 words recalled) and would be classified as demented on that basis alone. This is consistent with our general observation that most of the work of dementia discrimination is being performed by the three-item recall task, with the CDT acting to supplement and refine these judgements in less obvious cases. This is also consistent with the idea that three-item memory task is tapping into the first and most powerful factor in MMSE factor analyses. Additionally, we note that while the interpretation errors of mild CDT problems by naïve raters would decrease Mini-Cog sensitivity, it has the simultaneous effect of increasing specificity, resulting in no net change in overall correct detection. In an unselected population sample, we would predict that CDT-naïve raters using the Mini-Cog would have slightly lower sensitivity, slightly higher specificity, and very similar overall correct detection relative to more expert raters. In short, while naïve raters were less likely to identify mildly impaired CDTs as 'abnormal', we think that this would have minimal effects on overall Mini-Cog accuracy.

These results suggest that the Mini-Cog's 'minimalist' approach could be successfully used in settings where time is short, trained personnel are lacking, and/or language barriers exist. More lengthy and complex methods may be counter-productive, if physicians and other staff are unlikely to use them. The length and complexity of alternative instruments tends to defeat the goal of cost-effective dementia detection (Borson *et al.*, 2000), while the Mini-Cog speed and simplicity may promote it.

As is true for all first-stage screening tests, individuals who screen positive on the Mini-Cog cannot be presumed demented without additional evidence. Screening unselected elderly populations will result in increases in false-positive tests (lower specificity and diagnostic value) relative to the Mini-Cog developmental sample, which was enriched for probable cases. If we were to extrapolate the effects of using naïve raters in a population with substantially fewer impaired subjects, more false

negative and fewer false positives would be detected. This would have the advantage of decreasing the number of unnecessary diagnostic work-ups of non-demented older adults, but the disadvantage of reducing the detection of dementia in its earliest stages. However, we are encouraged by preliminary work comparing the Mini-Cog with the MMSE in a population based sample in Pennsylvania (the MoVIES sample: Ganguli *et al.*, 1993), which suggests that the Mini-Cog remains at least as sensitive and accurate as the MMSE, even when the CDT scoring system differs somewhat from that used here.

Full diagnostic dementia assessment of false-positive subjects is costly in time, money, and emotional distress. Previous studies have found that combining cognitive screening with informant history improves dementia detection over either method alone (Gurland *et al.*, 1995). Our approach, though different in detail, was similar in concept. For screening of elderly persons living alone without accessible informants, the Mini-Cog alone may be acceptable. However, a two-step screening approach, first using the Mini-Cog and then a brief informant interview, such as the short IQ-CODE (Jorm, 1994), warrants further population-based testing. If this method retains high sensitivity and minimizes false-positive screens, it may be the most cost-effective strategy for improving dementia detection and accurately selecting those in need of a complete diagnostic assessment.

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REFERENCES

- American Psychiatric Association. 1994. *Diagnostic and Statistical Manual of Mental Disorders*, 4th edn.
- Borson S, Brush M, Gil E, Scanlan JM, Vitaliano PP, Chen JC, Cashman J, Sta Maria MM, Barnhart R, Roques J. 1999. The **clock drawing test**: utility for dementia detection in multi-ethnic elders. *J. Gerontol. Med. Sci.* **54A**: M534–M540.
- Borson S, Scanlan JM, Brush M, Vitaliano PP, Dokmak A. 2000. The Mini-Cog: a cognitive 'vital signs' measure for dementia screening in multi-lingual elderly. *Int. J. Geriatr. Psychiatry.* **15**: 1021–1027.
- Braekhus A, Laake K, Engedal K. 1992. The Mini-Mental State Examination: identifying the most efficient variables for detecting cognitive impairment in the elderly. *J. Am. Geriatr. Soc.* **40**: 1139–1145.
- Brodsky H, Moore CM. 1997. The **Clock Drawing Test** for dementia of the Alzheimer's type: a comparison of **three scoring methods** in a memory disorders clinic. *Int. J. Geriatr. Psychiatry* **12**: 619–627.
- Bush C, Kozak J, Elmslie T. 1997. Screening for cognitive impairment in the elderly. *Can. Fam. Physician* **43**: 1763–1768.
- Dastoor DP, Schwartz G, Kurzman D. 1991. **Clock-drawing**. An assessment technique in dementia. *J. Clin. Exp. Geront.* **13**: 69–85.
- Death J, Douglas A, Kenny RA. 1993. Comparison of **clock drawing** with the Mini-Mental State Examination as a screening test in elderly acute hospital admissions. *Postgrad. Med. J.* **69**: 696–700.
- Fillenbaum GG, Wilkinson WE, Welsh KA, Mohs RC. 1994. Discrimination between stages of Alzheimer's disease with subsets of Mini-Mental State Examination items. An analysis of Consortium to Establish a Registry for Alzheimer's Disease data. *Arch. Neurol.* **51**: 916–921.
- Folstein MF, Folstein SE, McHugh PR. 1975. Mini Mental State: a practical method for grading the cognitive state of patients for the clinician. *J. Psychiatr. Res.* **12**: 189–198.
- Ganguli M, Belle S, Ratcliff G, Seaberg E, Huff FJ, van der Porten K, Kuller LH. 1993. Sensitivity and specificity for dementia of population-based criteria for cognitive impairment: the MoVIES project. *J. Gerontol.* **48**: M152–M161.
- Gurland BJ, Wilder DE, Chen J, Lantigua R, Mayeux R, Van Nostrand J. 1995. A flexible system of detection for Alzheimer's disease and related dementias. *Aging (Milano)* **7**: 165–172.
- Hill RD, Backman L. 1995. The relationship between the mini-mental state examination and cognitive functioning in normal elderly adults: a componential analysis. *Age Ageing* **24**: 440–446.
- Hughes CP, Berg L, Danziger WL, Coben LA, Masaki KH. 1982. A new clinical scale for the staging of dementia. *Br. J. Psychiatry* **140**: 566–572.
- Jorm AF. 1994. A short form of the Informant Questionnaire on Cognitive Decline in the Elderly

- (IQCODE): development and cross-validation. *Psychol. Med.* **24**: 145–153.
- Jorm AF, Scott R, Cullen JS, MacKinnon AJ. 1991. Performance of the Informant Questionnaire on Cognitive Decline in the Elderly (IQCODE) as a screening test for dementia. *Psychol. Med.* **21**: 785–790.
- Juby A. 1999. Correlation between the Folstein Mini-Mental State Examination and **three methods of clock drawing scoring**. *J. Geriatr. Psychiatry Neurol.* **12**: 87–91.
- Klein L, Roca R, McArthur J, Vogelsang G, Klein GB, Kirby SM, Folstein M. 1985. Diagnosing dementia: univariate and multivariate analyses of the mental status examination. *J. Am. Geriatr. Soc.* **33**: 483–488.
- Lam LC, Chiu HF, Ng KO, Chan C, Chan WF, Li SW, Wong M. 1998. **Clock-face drawing**, reading and setting tests in the screening of dementia in **Chinese** elderly adults. *J. Gerontol. B Psychol. Sci. Soc. Sci.* **53**: 353–357.
- Magaziner J, Bassett SS, Hebel JR. 1987. Predicting performance on the Mini-Mental State Examination. Use of age- and education-specific equations. *J. Am. Geriatr. Soc.* **35**: 996–1000.
- McKhann G, Drachman D, Folstein MF, Katzman R, Price D, Stadlan EM. 1984. Clinical diagnosis of Alzheimer's disease: report of the NINCDS-ADRDA work group under the auspices of the Department of Health and Human Services Task Force on Alzheimer's Disease. *Neurology* **34**: 939–944.
- Roca RP. 1987. Bedside cognitive examination. Usefulness in detecting delirium and dementia. *Psychosomatics* **28**: 71–76.
- Shulman K, Shedletsky R, Silver I. 1986. The challenge of time: **clock-drawing** and cognitive function in the elderly. *Int. J. Geriatr. Psychiatry* **1**: 135–140.
- Teng EL, Hasegawa K, Homma A, Imai Y, Larson E, Graves A, Sugimoto K, Yamaguchi T, Sasaki H, Chiu D. 1994. The Cognitive Abilities Screening Instrument (CASI): a practical test for cross-cultural epidemiological studies of dementia. *Int. Psychogeriatr.* **6**: 45–58.
- Tuokko H, Hadjistavropoulos T, Miller JA, Beattie BL. 1992. The **Clock Test**: a sensitive measure to differentiate normal elderly from those with Alzheimer disease. *J. Am. Geriatr. Soc.* **40**: 584.
- Watson YI, Arfken CL, Birge SJ. 1993. **Clock completion**: an objective screening test for dementia. *J. Am. Geriatr. Soc.* **41**: 1235–1240.
- Yuspeh RL, derploeg RD, Kershaw DA. 1998. Validity of a semantically cued recall procedure for the mini-mental state examination. *Neuropsychiatry Neuropsychol. Behav. Neurol.* **11**: 207–211.