



Introduction

 Depression is one of the most prevalent non-cognitiv correlates of AD, with prevalence estimates mos commonly 30-50%.

• The effect of depression on cognition in AD has no been consistently demonstrated; however, evidenc suggests reduced performance on measures processing speed and attention in depressed patient with AD.

 Cognitive reserve is thought to delay the onset an slow the progression of cognitive impairments in AD. B the same reasoning, cognitive reserve may protect against the effects of depression on cognition patients with AD.

 Educational and occupational attainment, as well a higher scores on mental ability tests, are correlated wit cognitive reserve. These same variables ar significantly related to better functioning and copin after brain injury or disease.

• The present study aimed to examine the effects cognitive reserve on the relationship betwee depressive performance symptoms and neuropsychological tests in patients with probable AD. Hypothesis: Cognitive reserve moderates relationship between depressive symptoms and performance on neuropsychological tests in patients with probable AD.

Participants

 Prospectively collected longitudinal data was obtained from the Baylor College of Medicine Alzheimer's Disease and Memory Disorders Center.

 Inclusion criteria included meeting NINCDS-ADRDA criteria for probable AD and MMSE score > 14 so as to only include patients who could adequately report their depressive symptoms on the GDS (n = 520).

• The final sample was 67% female and the mean age was 74.63 years (SD = 7.97).

Cognitive Reserve as a Moderator for the Relationship between Depression and Cognitive Functioning in Alzheimer's Disease

Whitney N. Havins^a, Kristina A. Agbayani^a, Paul J. Massman, Ph.D.^{ab} & Rachelle S. Doody, M.D., Ph.D.^b

^a University of Houston, ^b Baylor College of Medicine

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Kesults

• Cognitive reserve was significantly related to performance on all of the measures administered. Depressive symptoms were neuropsychological significantly related to processing speed and phonemic fluency. • Cognitive reserve demonstrated incremental validity, over and above depressive symptoms, in predicting cognitive test performance. • Cognitive reserve accounted for significant variance over and above depressive symptoms for processing speed (12%), verbal episodic memory (7%), visual episodic memory (5%), phonemic fluency (20%), and semantic fluency (5%); but not for overall level of cognitive dysfunction (ADAS). In multiple regression analyses, the interaction between GDS and AMNART scores approached significance in predicting performance on VSAT time. Thus, additional analyses on this variable were conducted after grouping the subjects by AD severity.

• Regression analyses conducted on patients with mild AD showed a significant interaction between GDS and AMNART scores when predicting VSAT time (p < .05). This suggests that cognitive reserve moderates the relationship between depressive symptoms and processing speed in these patients (Figure 1 & 2).

all selected tests administered (see

as used to predict performance on MNART estimated IQ scores. patients with mild AD (MMSE \geq 20)

derated the relationship between is subset of the sample.

ctions Measured

VE FUNCTION MEASURED

e Reserve

ive symptoms

nental status

e dysfunction in AD

memory (Verbal and Visual)

ic Fluency

c Fluency and psychomotor speed Figure 1. Effect of Cognitive Reserve and Depressive Symptoms on VSAT Time (Attention/Psychomotor Speed—Lower Values Better, N = 299).



Figure 2. Standardized Regression Coefficients for the Effect of Cognitive Reserve, Depressive Symptoms, and the Interaction in Predicting **Psychomotor Speed (N = 299)**

Interaction

 Results from this study showed that depressive symptomology is not a reliable predictor of performance on neuropsychological testing in patients with AD. • However, cognitive reserve (either independently or by association with educational and occupational achievement) consistently predicts and accounts for significant variance in cognitive ability. Cognitive reserve moderated the relationship between depressive symptoms and psychomotor speed in patients with mild AD. Patients with high cognitive reserve and fewest depressive symptoms performed best on a speeded task. The attention/psychomotor speed of AD patients with high cognitive reserve was more negatively affected by depressive symptoms than that of patients with lower cognitive reserve.



Cognitive Reserve (AMNART IQ)

Symptoms (GDS)

Cognitive Reserve

 β = .-.48, *p* < .001

Psychomotor Speed

Depressive Symptoms

 β = -1.01, *p* = .078

Conclusions

 $\beta = 1.10, p < .05$