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Semantic feature analysis for treatment of anomia in early Alzheimer's disease: Two cases studies

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INTRODUCTION

Alzheimer's disease (AD) is one of the most common neurodegenerative diseases. In the early stages of the disease, a semantic memory deterioration can be observed, manifesting itself through lexico-semantic difficulties as anomia, semantic paraphasia and circumlocutions. Semantic Feature Analysis (SFA), proposed by [1] and developed by [2-3], aims to reduce anomia in patients with aphasia by reinforcing lexico-semantic network. However, studies regarding the efficiency of SFA in AD are scarce [4-5]. The aim of this study was to investigate the effect of SFA on anomia in early AD.



PROCEDURE Naming abilities of participants were assessed by a naming task of 100 pictures during the pretest phase. Then, 15 concepts among the failed pictures were trained with SFA chart (figure 1) for eight weeks. Twice a week, participants attended a 60-minutes session. Finally, the naming abilities were reassessed in the posttest and follow-up phases.

RESULTS

To compare naming task performances at different assessment phases, the Q of Cochran statistic was used. Z-score were used for two by two comparisons.

Participant MS. Results showed a significant improvement (Q(2) = 19.5; p < .001) in naming performances, particularly between pretest and posttest phases (z = 3.34; p <.001 *) (figure 2). Moreover, the qualitative analysis of lexical errors (figure 3) showed a decrease of the non-response rate. Indeed, MS produced more semantic paraphasias, as well as more circumlocutions.





<u>Participant MV</u>. No significant improvement in naming was observed (Q(2) = 1.28; p > .05) (figure 4). However, the qualitative analysis of lexical errors showed a change in lexical production (figure 5). Indeed, the non-response rate decreased and more semantic paraphasias and circumlocutions were observed. Non-respon



DISCUSSION AND CONCLUSIONS

In this study, we explored the benefits of SFA in two case studies, MS and MV. The method was efficient only for MS, showing a significant improvement in naming as well as a sustained benefit in the follow-up. We also observed a semantic reorganization, with fewer non-responses and an increase in lexical productions. In contrast, MV's naming performances did not significantly change. This lack of response could be partly explained by a more severe general cognitive and semantic decline. While we observed no improvement in MV, there was an increase in lexical productions, albeit erroneous in posttest phase. In conclusion, the SFA-based treatment of anomia yielded significant positive evolutions in one of our AD participants, reinforcing her lexical-semantic network, given that the semantic deterioration was not too severe. Our initial findings provide evidence-based recommendations for managing anomia in AD, though more research is needed to support our preliminary results.

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