Relationship between hippocampal subfields and category cued recall in **AD and PDD: a multimodal MRI study**



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OBJECTIVE: We investigate how changes in whole hippocampus and hippocampal subfields relate to memory recall in different forms of dementia, as Alzheimer's Disease (AD) and Parkinson's Disease with Dementia (PDD)

MATERIALS AND METHODS: Twenty-two AD subjects, 18 PDD and 17 healthy controls were recruited for a study protocol in which a multimodal 3T-MRI hippocampal evaluation (whole-brain T1-weighted and Diffusion Tensor Imaging [DTI]) was combined with a hippocampal targeted neuropsychological assessment (Free and Cued Selective Reminding Test [FCSRT]). Macro- and micro-structural features (volume; shape; mean diffusivity [MD]; fractional anisotropy [FA]) of bilateral hippocampi (whole and subfields) were obtained. Correlations between MRI-derived parameters and neuropsychological evaluations were performed

RESULTS: Compared to controls, AD showed a reduction in total hippocampal volume and in almost all subfields, with a MD increase in the same regions, whereas PDD displayed a volume loss, less severe than AD, more evident in the CA2-3 and presubiculum subfields, without a clear consensual MD increase. In the comparison between AD and PDD, the multimodal analysis allowed us to identify that subiculum, CA1 and CA4-DG were the subfields differently involved in these diseases. Interestingly the same subregions correlated with immediate and delayed total recall items (ITR and DTR) of FCSRT, thus suggesting that these subfields play a role in the total cued recall



Hippocampal Shape analysis comparison between groups



p-values

Correlation analysis in AD group between hippocampal shape and FCSRT scores







Subicular Complex



DTR



CONCLUSIONS: Our study provides compelling new evidence that hippocampal subregions had different vulnerability to damage related to different forms of dementia. Moreover, the combination of the *in vivo* analysis of hippocampal subfields with the FCSRT paradigm provided important insights into whether changes within specific hippocampal subfields are related to the different mnesic profile in AD and PDD patients

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