# Visual pathway abnormalities in dementia with Lewy bodies: an Optical Coherence Tomography (OCT) and 18F-FDG-PET/MRI study

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### Background

The visual system may be involved in several core features of dementia with Lewy bodies (DLB), however, very few data on retinal and visual system abnormalities are available. Aim: To identify structural and metabolic features alongside the visual pathway (i.e. from retina to primary/secondary visual cortex) that may be specific of DLB.

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## Methods

<u>Study 1</u>: 35 DLB patients and 30 healthy subjects (HS) underwent a Spectral Domain retinal Optical Coherence Tomography (OCT) scan; peripapillary RNFL (pRNFL) and macular layers thicknesses and volumes were compared. Exclusion criteria for all subjects were presence of retinopathy, severe glaucoma and age-related macular degeneration. DLB patients underwent clinical interview, neurological and neuropsychological evaluations.

<u>Study 2</u>: 31 DLB patients and 18 control subjects underwent 18F-FDG-PET/MRI scan. Cortical thickness (Cth), subcortical volumes and 18F-FDG SUVr of visual system structures were compared.

<u>Study 3</u>: data from 16 DLB patients with both OCT and 18F-FDG-PET/MRI were combined to identify associations between retinal thickness and brain metabolism of the visual pathway.

#### Results

<u>Study 1</u>: pRNFL thickness was not significantly different in DLB patients and HS; as for macular layers, parafoveal ganglion cells and inner plexiform layer (pfGCIPL) was significantly thinner in DLB patients Fig. 1: Comparison of pRNFL values between DLB patients and healthy subjects





(p=0,03), mostly in nasal and temporal quadrant (p=0,02), as well as the inner nuclear layer (INL) inner temporal quadrant (p=0,003). The thickness of pRNFL temporo-superior sector was associated with disease duration (Rho=-0,4, p=0,01), while visuo-spatial abilities were associated with the thickness of pRNFL temporal sector and papillomacular bundle (QPST "opening/closure" item: Rho=0,5, p=0,004 and Rho=0,4, p=0,01, respectively; QPST "closing in" item: Rho=0,5, p=0,002 and Rho=0,5, p=0,001, respectively).

<u>Study 2</u>: DLB patients had significantly thinner Cth in secondary visual areas, i.e. right precuneus (p=0,003) and bilateral fusiform gyrus (p=0,02, both); and lower SUVr in parieto-temporo-occipital regions.

<u>Study 3</u>: pfGCIPL thickness and volumes of macular GCIPL (GCIPLv), RNFL (RNFLv) and INL (INLv) had a negative association with level of glucose metabolism in the fusiform gyrus (FG) and pulvinar (FG: pfGCIPL thickness r=-0,6, p=0,03; GCIPLv r=-0,6, p=0,03; RNFLv r=-0,7, p=0,003. Pulvinar: INLv r=-0,6, p=0,01; GCIPLv r=-0,5, p=0,04).

## Conclusion



in DLB patients parafoveal macular GCIPL and Cth in secondary visual areas are thinner than in control subjects and glucose metabolism lower in temporo-parieto-occipital cortical regions. Thicknesses of macular layers in DLB group were negatively associated to secondary visual cortex metabolism, suggesting a relative preservation of synaptic activity in secondary visual cortex in response to degraded afferent visual stimuli.

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