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### **OBJECTIVE**

Although that other non-motor, prodromal signs of Parkinson's disease (PD) are well described, only little is known about the cognitive profile in the prodromal phase.

This study investigates the cognitive profile in people with REM-sleep Behavior Disorder (RBD) and hyposmia, in a prodromal PD cohort. Knowledge of these alterations in prodromal PD is still limited<sup>3</sup>.

# **BACKGROUND**

PD PRODROMAL PERIOD can start

20 years

before onset of motor parkinsonism

**RBD & OLFACTORY** 

DYSFUNCTION

= RISK FACTORS

for

alpha-synucleinopathies,

such as PD

75%
Of people with
RBD
Progress to PD
within 10 years

Studies suggested that changes to EXECUTIVE FUNCTION

in particular, may be considered as **non-motor marker** of prodromal PD

Prospective longitudinal studies are needed to assess the emergence of cognitive symptoms over time and to define prodromal non-motor symptoms.

**Descriptive statistics** P-Values Significance Variable pRBD nRBD pRBD vs. (n = 95)SD Mean SD 63 / 32 63 / 32 p = 1.000 Gender, M / F 62.95 62.92 8.46 p = 0.245Age, in years 12.90 p = 0.80025.28 2.49 p < 0.001

Table 1. Demographic and clinical data for pRBD and nRBD: SD: Standard Deviation; M: Male; F: Female; n = sample size; MoCA: Montreal Cognitive Assessment; \* Significant at the 5% level (2-tailed). \*\* Significant at the Bonferroniadjusted 5% level (p-value <-0.05/4)

After multiple testing correction, we observed that MoCA subitems assessing visuo-constructive

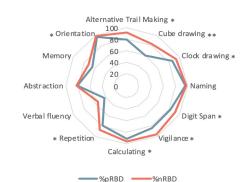
abilities were significantly lower in pRBD + hyposmia compared to the control group without RBD-hyposmia.

Furthermore, pRBD showed nominally lower scores in items assessing executive funtions, visual-spatial functions and orientation.

### PRELIMINARY RESULTS

We observed lower MoCA total scores (p < 0.001) in the pRBD group, suggestive for a lower global cognitive profile in the pRBD and hyposmia group compared to the matched control group (Table 1).

MoCA subitems comparison pRBD vs. nRBD sting correction, we observed



## **METHODS**

pRBD & Hyposmia N=95 No RBD & No Hyposmia N=95

Natched for age and gender No sign. difference for education

MoCA
Global cognition

Cross-sectional study of 190 participants from the Luxembourg RBD Study.

Participants were assigned to the probable RBD and hyposmia (pRBD; n=95) or no RBD and no hyposmia (nRBD; n=95) groups based on:

- RBD Screening Questionnaire (RBDSQ)<sup>5</sup>
- Olfactory test (B-SIT/Sniffin'Stick)<sup>6,7</sup>.

The Montreal Cognitive Assessment (MoCA)<sup>8</sup> was applied to assess global cognition. Broader neuropsychological assessments are foreseen in the coming weeks to validate the preliminary results in the full sample of the Luxembourgish RBD study. Excluded were people with PD, as well as other neurological diseases and severe psychiatric disorders

# PRELIMINARY RESULTS ON THE COGNITIVE PROFILE IN PRODROMAL PARKINSON'S DISEASE - A PROSPECTIVE STUDY

### OUTLOOK

Future research will seek to validate these results with broader neuropsychological assessments, polysomnography and increased sample size in the Luxembourgish RBD study. Furthermore, longitudinal analysis will evaluate conversion rates to alpha-synucleinopathies.

Poster in PDF:



For more information about the Luxembourgish RBD Study:



results show that:

**SUMMARY** 

 $\searrow$  Global cognition in pRBD+hyposmia compared to the control group. Besides this, pRBD performed worse in visuospatial functions.

In line with the literature on prodromal PD3, our

We observed tendencies for impaired executive functions in pRBD. Given, that these observations are based only on sub-items of a screening tool, we need to be careful with the interpretation.



