

## Gait Analysis and Pattern Cognition in Parkinson's Disease

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**Background:** The relationship between cognition and gait in Parkinson's disease (PD) has received increased attention, however the specific connection between gait features and cognition is still not fully understood.

**Aim:** To find association between gait parameters and specific cognitive profiles in patients with PD.

**Methods:** Using motion analysis system, forty three PD patients during gait in normal conditions at on state were studied. We evaluated the following gait parameters: 1) step length; 2) stance phase; 3) swing phase; 4) single support/double support time ratio; 5) cadence; 6) velocity; 7) step length variability; 8) swing time variability. In order to reduce the larger number of gait variables to a smaller number of elements, a factor analysis was performed. Furthermore, we assessed all patients with an extensive neuropsychological battery and correlated the composite scores of three main cognitive domains, namely episodic memory, executive and visuospatial domains with the gait factors scores.

**Results:** Factor analysis revealed two independent factors, namely "pace" and "stability". The "pace" factor was not correlated with cognitive or clinical variables. The "stability" factor was strongly and directly correlated with visuospatial domain.

**Conclusions:** Visuospatial impairment was strongly associated with the development of instability and the progression of disease.

## Genetic Risk Factors in Neurodegenerative Diseases

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**Background:** The risk of developing Alzheimer's disease (AD) and frontotemporal dementia (FTD) could be associated to SNPs in specific genes.

**Aim:** To define a specific profile of risk for AD and FTD.

**Population and methods:** SNPs of APOE, Cyp46A1, ABCA1, PRNP, TOMM40, GAB2, NOS3 genes analyzed in 322 AD patients, 102 FTD patients and 366 controls. Statistics performed by  $\chi^2$  test.

**Results:** The APOE  $\epsilon 4$  allele frequency was significantly different ( $p=0.000$ ) in AD (22.1%) and FTD (18.6%) subjects than controls (7.8%) with an OR of 3.36 and 2.71, respectively. Genotypic frequency of the GAB2 gene in FTD patients was different than controls ( $p<0.006$ ), with a frequency of TT genotype of 10% in subjects FTD vs. 2.8% in controls, and a T allele frequency was of 27% vs. 19% in controls ( $p=0.02$ ), with an OR of 1.52. Patients with TT and  $\epsilon 4$  had an increased risk of developing FTD (OR7.28).

**Conclusions:** The association between APOE  $\epsilon 4$  allele with AD and FTD is once again confirmed. T allele homozygosity of the GAB2 gene associated to  $\epsilon 4$  increases the risk of developing FTD but further studies are needed to understand the role of this gene in the biochemical mechanisms underlying neurodegeneration.

## Two Cases of Early Onset FTLT Due to the Chromosome 9 Hexanucleotide Repeats

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## **PROCEEDINGS**

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