

MEETING ABSTRACT

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# Early involvement of sympathetic cardiac nerve endings in a patient with rem sleep behaviour disorders

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## Background

REM sleep behaviour disorder (RBD) is a parasomnia characterized by suppression of muscular atonia and abnormal, often violent, motor behaviours during REM sleep. RBD can be either idiopathic or associated with degenerative disorders, such as dementia with Lewy bodies (LBD), multiple system atrophy (MSA), progressive supranuclear paralysis (PSP) or Parkinson's disease (PD). Myocardial <sup>123</sup>I-Metaiodobenzylguanidine (MIBG) enables the assessment of postganglionic sympathetic cardiac nerve terminals. Reduced myocardial MIBG uptake of the tracer suggests cardiac sympathetic denervation. MIBG uptake allows a differentiation between Lewy body disease and tauopathies and other parkinsonisms. Indeed cardiac innervation is impaired in nearly all patients with PD and LBD, while it is preserved in patients with parkinsonisms, such as MSA and PSP. Recently, decreased MIBG uptake has been also described in patients with RBD.

## Materials and methods

A 72-year-old woman reported a history characterized by frequent nocturnal nightmares, vivid dreams, strong shaking of her limbs, laughing, talking, screaming and frequent falls from the bed. There was no family history of neurological disorders or RBD. When she was first admitted to our clinic (May 2008) her neurological examination showed only a slight facial hypomimia. All cognitive tests were normal, the MMSE showed a score of 28/30 corrected (n.v. > 24/30). Routine biochemistry (including serum glucose), haematological tests, electrocardiogram, echocardiography, Holter blood pressure

monitoring, brain magnetic resonance imaging were all normal. A polysomnographic examination showed the characteristic features of RBD. In order to investigate the nigrostriatal system integrity, we performed a cerebral SPECT with <sup>123</sup>I-FP-CIT that resulted normal. The cardiac MIBG scintigraphy showed, however, a markedly reduced cardiac uptake.

## Discussion and conclusions

RBD may represent a prodromic phase of neurodegenerative diseases, such as PD, MSA or LBD. In our case, the RBD was the only relevant clinical manifestation in absence of signs of extrapyramidal dysfunction. The DAT-scan of the patient was normal, while the myocardial MIBG scintigraphy showed a marked impairment of sympathetic innervation. This is the first evidence of a patient with RBD with cardiac sympathetic denervation and a normal DAT-scan, demonstrating that cardiac sympathetic nerve terminals may be impaired in absence of detectable damage in nigrostriatal the cardiac innervation may precede neuronal cell loss in the dopaminergic nigrostriatal system.

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