Dear Sir,

Narcolepsy (NC) is a sporadic hypersomnia (prevalence 1:2,000), characterized by excessive daytime sleepiness (EDS) and sleep attacks, typically associated with cataplexy and other REM-sleep related phenomena such as sleep paralysis and hallucinations [1]. In human NC with cataplexy, an association with specific HLA haplotype (DR DQB1 0602) and a deficiency in hypocretin (orexin) peptide in cerebrospinal fluid (CSF) are almost constant findings. The pathogenesis of human NC is still unknown. Autoimmune and neurodegenerative processes of hypothalamic structures have been discussed with more solid evidence for the first theory [2–4]. Most of narcoleptic symptoms develop early after the onset of the disease and usually do not worsen with the progression of the neurodegeneration [2]. Moreover, the absence of ubiquitinated inclusions (cardinal neuropathological finding of most neurodegenerative diseases) in narcoleptic patients argues against the neurodegenerative hypothesis in NC [3]. An autoimmune process involving the hypocretin neurons of lateral hypothalamus has been documented with post-mortem [4].

There are few reports in the literature showing the occurrence of NC-like symptoms in patients already affected by Parkinson’s disease (PD) and other neurodegenerative diseases [2, 5–7]. Hypocretin neuronal loss has been documented in subjects with advanced PD and Alzheimer’s disease (AD) [6, 7]. A possible relationship between neurodegenerative disorders, especially PD as a possible cause of typical NC with hypocretin deficiency and HLA haplotype, and NC-like phenotype is still under discussion.

Herein, we describe 2 patients with a long history of NC who developed PD and AD, respectively.

**Case Reports**

**Case 1**

A 69-year-old Caucasian male, suffering from NC since the age of 17, presented to us at the age of 64 years (in 2006). His history included EDS with sleep attacks, hallucinations, frequent cataplexy and occasional sleep paralysis. Patient’s psychomotor development was normal and clinical history unremarkable; in particular, he denied infectious or inflammatory cerebrospinal fluid (CSF) were already tried without efficacy. Conversely, cataplexy was partially improved by anticholinergic agents – biperiden, and modafinil and imipramine and presented a resting tremor, micrographia, and slight bradykinesia of the right hand. Hyposmia was documented with olfactory tests. Total Unified Parkinson’s Disease Rating Scale (UPDRS III) score was 13 (mental activities: 0, activities of daily living: 6, motor: 7; UPDRS III range is 0–176). The brain MRI was normal. Before our evaluation, the tremor was diagnosed as essential (he had a family history of essential tremor) and specific treatments (beta-blockers, primidone and anticholinergic agents – biperiden) were already tried without efficacy. Conversely, cataplexy was partially improved by the anticholinergic treatment (biperiden), while other narcoleptic symptoms remained unchanged. Because of the limited response to the therapy, the patient was started on L-dopa-benserazide (until the final dose of 1,000 mg/day), which ameliorated all symptoms, with a decrease of 70% in motor UPDRS score (total score: 3, mental activities: 0, activities of daily living: 1, motor: 2). Thus, according to the standard diagnostic criteria, the diagnosis of PD was done.
Soon after the introduction of L-DOPA, symptoms of NC also drastically improved, in particular cataplexy, which remitted completely with the exception of episodes of short duration of eyelid ptosis on emotional triggers; EDS improved as well (Ewpoor Sleepiness Scale 11/24). The above-described NC status remained unchanged until the last clinical assessment (at the age of 68, in 2010).

**Case 2**

A 71-year-old Caucasian male suffered from NC symptoms since his adolescence. He always claimed severe EDS with recurrent sleep attacks. Frequent cataplectic episodes were also experienced since adolescence, usually triggered by positive emotions (mainly laughing). Insomnia (fragmented sleep) and hypnopompic hallucinations but no sleep paralysis were reported. Because of the severe disability, the patient was exempted from his military duty and preferred working at home as a painter, adjusting his daily activities to his sleep/wake regulation. Some considerations regarding the appearance of neurodegenerative disorders in subjects already affected by NC [3, 7]. Especially as far as AD is concerned, the reports in the literature document postmortem diagnosis of AD. Thus, our patient is the first one reported suffering from NC in whom diagnosis of AD according to the international criteria was done when still alive, receiving consequently the appropriate therapy.

The prevalence of PD and AD at the age and in the geographic area of the 2 patients described here, is estimated at 9.5/1,000 for PD and 48/1,000 for AD, respectively [8], while the prevalence of NC is around 0.5/1,000 [1]. Thus, the probability for NC patients to present comorbid PD and AD can be estimated at about 4.75/1,000,000 and 24/1,000,000 respectively. Although infrequent, these proportions are not extremely rare (especially when considering referral centers for NC), and may suggest a rare but not negligible comorbidity. A recent neuropsychological study on 12 NC patients showed a similar prevalence of AD among them compared to the prevalence of AD in the general population [7]. Although no available evidence of a predisposition of NC to neurodegenerative diseases (PD and AD in particular) exists, given that both dopaminergic and cholinergic systems are implicated in the sleep/wake regulation, some considerations regarding their role, their dysregulation and also their complex interactions with the hypocretin neuronal networks might be discussed.

Hypocretin deficiency due to hypocretin cell loss in lateral hypothalamus is the

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**Table 1. Clinical features before and after neurodegenerative disorder development**

<table>
<thead>
<tr>
<th>Clinical feature</th>
<th>PD patient before PD development</th>
<th>4 years after PD development</th>
<th>AD patient before AD development</th>
<th>4 years after AD development</th>
<th>6 years after AD development</th>
</tr>
</thead>
<tbody>
<tr>
<td>EDS</td>
<td>severe (ESS 20)</td>
<td>moderate (ESS 11)</td>
<td>severe (ESS 16–18)</td>
<td>slight (ESS 8)</td>
<td>n.a.</td>
</tr>
<tr>
<td>Mean SL, min</td>
<td>6.6 (treated)</td>
<td>n.a.</td>
<td>1.9</td>
<td>4.5</td>
<td>1</td>
</tr>
<tr>
<td>Cataplexy</td>
<td>severe, daily</td>
<td>rare, only eyelid</td>
<td>rare, only eyelid</td>
<td>no</td>
<td>no</td>
</tr>
<tr>
<td>SOREM</td>
<td>4</td>
<td>n.a.</td>
<td>2</td>
<td>2</td>
<td>5</td>
</tr>
<tr>
<td>Sleep paralysis</td>
<td>yes</td>
<td>rare</td>
<td>no</td>
<td>no</td>
<td>no</td>
</tr>
<tr>
<td>Hallucinations</td>
<td>yes</td>
<td>rare</td>
<td>frequent</td>
<td>rare</td>
<td>frequent</td>
</tr>
<tr>
<td>CSF Hyp-1</td>
<td>positive</td>
<td>–</td>
<td>undetectable</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>HLA DQB1*0602</td>
<td>positive</td>
<td>–</td>
<td>positive</td>
<td>–</td>
<td>–</td>
</tr>
</tbody>
</table>

ESS = Epworth Sleepiness Scale; Hyp-1 = hypocretin; n.a. = not available.

Discussion

While the development of NC-like syndromes secondary to neurodegenerative disorders (mostly PD) is well described in the literature, there are only isolated reports regarding the appearance of neurodegenerative disorders in subjects already affected by NC [3, 7]. Especially as far as AD is concerned, the reports in the literature document postmortem diagnosis of AD. Thus, our patient is the first one reported suffering from NC in whom diagnosis of AD according to the international criteria was done when still alive, receiving consequently the appropriate therapy.
hallmark neuropathological damage in NC. A partial hypocretin deficiency seems to be also common in PD and was recently shown in an AD postmortem study as well [6, 7]. In PD, an association with narcoleptic-like symptoms, such as dreaming-like phenomena while awake, hypnagogic hallucinations and REM behavior disorder, which often precede the appearance of motor symptoms of PD, is not a rare phenomenon, suggesting a progressive hypocretin cell loss of degenerative origin.

In the first patient described here, the PD onset was not followed by a worsening of the narcoleptic symptoms. Interestingly, the dopaminergic treatment improved EDS and cataplexy allowing the withdrawal of the NC medication. This finding is in contrast with the common knowledge regarding dopaminergic treatments, which not only do not enhance vigilance, but also may induce sleep attacks both in PD patients and in healthy controls [9–11].

Sleep-wake pattern abnormalities in PD patients could depend on both hypocretin deficiency and dopaminergic dysfunction. In fact, the dopaminergic neurons play an important role in promoting wakefulness. An impairment of the dopaminergic system has also been suggested in NC, for instance D2-receptor binding was elevated in NC and positively correlated with the frequency of cataplectic and sleep attacks. Furthermore, an impairment of the dopamine-mediated reward system has also been described [12–14]. This observation suggests a complex interaction between dopaminergic and hypocretin neuronal networks, which needs further investigations.

Acetylcholine plays an important role in promoting vigilance toward a diffuse cortical activation. A cholinergic deficit is a hallmark of AD. Dysfunction in orexin–acetylcholine interactions may play a role in the arousal and attention deficits seen in neurodegenerative conditions, in drug addiction and in age-related cognitive decline as well [15]. Moreover, data from canine NC studies suggested that a primary deficit in orexin signaling might contribute to postsynaptic degeneration and affect Ach-dependent cognitive functions [16]. Under this prospective, it would be expected that the occurrence of a neurodegenerative disorder could worsen at least some of the NC symptoms, especially EDS. Conversely, in both of our patients the appearance of neurodegenerative process was followed by a significant improvement of all narcoleptic symptoms, including EDS and cataplexy. EDS’s paradoxical improvement, as already reported [17], may be due to the overall sleep improvement and sleep-wake cycle regulation following cholinergic treatment. Interestingly, cataplexy is also mediated by the cholinergic system, and usually responds to anti-cholinergic drugs, while in our case, cataplexy was improved by an enhancement of the cholinergic system. This observation together with the overall amelioration of the narcoleptic symptoms after the clinical onset of the neurodegenerative diseases confirms the complexity of the interactions between the hypocretin/dopaminergic and cholinergic systems.

In this scenario, a long-term follow-up of narcoleptic patients becomes extremely relevant to better understand the above-mentioned complex neuronal network interactions and to improve the therapeutic approach in narcoleptic patients with neurodegenerative comorbidity.

References