Piribedil is a dopamine agonist used to treat Parkinson’s disease, but its efficacy is less well evaluated than that of other dopamine agonists (1,2). It is also authorised in France as a “vasodilator” to treat certain types of chronic cognitive and neurosensory deficits in elderly patients, intermittent claudication caused by chronic peripheral artery occlusive disease, and in ophthalmology, although its efficacy in these indications has not been demonstrated (2-5).

Piribedil exposes patients to the adverse effects of dopamine agonists, including sudden sleep attacks, most of which have been reported in patients with Parkinson’s disease (a)(1.6). Does this adverse effect also occur when piribedil is used for other disorders, by patients without Parkinson’s disease? A case series published in 2011 provides some answers to this question.

Sudden sleep attacks, from the first few days of treatment. A group from the Lyon Regional Pharmacovigilance Centre searched the French national pharmacovigilance database for reports of sleep attacks attributed to piribedil in patients without Parkinson’s disease, recorded between January 1988 and December 2008 (7).

According to the definition adopted in this study, sleep attacks are “events of overwhelming sleepiness that occur without warning or with a prodrome sufficiently short or overpowering to prevent protective measures” (7).

The French pharmacovigilance database contained 7 reports of sleep attacks that met this definition in patients without Parkinson’s disease who were taking piribedil. The average age of these 7 patients was 69 years. The daily dose ranged from 20 mg to 150 mg (median: 50 mg). Six patients were using piribedil for a variety of vascular disorders, and the other patient was taking it for essential tremor. In 6 cases, piribedil was the only drug implicated (7).

In 5 cases, the sleep attacks began within 3 days of taking the first dose. In one woman who had taken piribedil for 10 years with 2 interruptions, the sleep attacks stopped after each discontinuation and recurred each time the drug was reintroduced (7).

Road traffic accidents, falls. Four patients experienced sleep attacks while working or driving. One 56-year-old man was admitted to hospital following sleep attacks that occurred two hours after his second dose of piribedil. One 75-year-old man had an initial sleep attack while he was driving, one hour after taking his first dose of piribedil. He felt the need to stop for a 10-minute nap. He had two more attacks after resuming his journey, the second of which caused a road traffic accident. One 86-year-old man taking piribedil for peripheral vascular disorders had two falls caused by sleep attacks. The patients described these sleep attacks as intense and sudden. In all cases, the sleep disorders resolved after discontinuation of piribedil (7).

A direct role for piribedil. Sleep attacks are a known adverse effect of dopamine agonists. As most of the data came from patients with Parkinson’s disease, uncertainty remained over the role of the disease in these sleep disorders (6). These case reports, in patients taking piribedil but who did not have Parkinson’s disease, are consistent with this drug having a direct role in sleep attacks, and probably a similar role to that of other dopamine agonists.

In practice: piribedil has little value. Piribedil has not been shown to be an effective treatment for vascular disorders. It is not the first-choice dopamine agonist in Parkinson’s disease, and its value is unclear (1,3,8). Unnecessary exposure of patients to the adverse effects of piribedil is not justified. It is better to use drugs with demonstrated efficacy instead.